

Electronic Structure of Impurities and Vortices in Short Coherence Length Superconductors

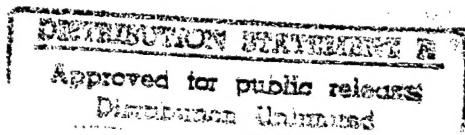
Annual Report

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PROGRAM GOALS

This grant is for a twenty-seven month investigation of the influence of impurities and vortices on the nearby electronic structure in a short coherence length superconductor. The primary goal is to develop self-consistent calculations of the electronic structure near various defects, focusing on the local density of states (LDOS) measurable by an STM. These calculations are to be compared with analytic results in various limits. Pairing functions of *d*-wave and *s*-wave symmetry are to be considered. Results for the local electronic structure are expected to be of great utility in understanding macroscopic measurements such as planar tunneling spectra, critical currents and the magnetic penetration depth, but the application of theoretical results to these problems is currently not funded.

PROGRESS

Over the past year we have developed a new Green's function technique, which we call the Koster-Slater technique, for calculating the local electronic structure in superconductors. This technique allows us to calculate the changes in the order parameter self consistently and is applicable to weak or strong-coupling superconductors with arbitrary order parameters. In a practical numerical calculation the Koster-Slater technique allows for essentially arbitrarily fine energy resolution in the calculated tunneling spectra, and requires approximately one hour to run for a high-temperature superconductor on a high-end desktop workstation. In contrast, the Bogoliubov-de Gennes equations have lousy energy resolution for realistic superconducting energy gaps, and we estimate would require six months to run for a single spectrum (if there were a computer with one million MB of memory).

The initial application of this technique was to an isotropically gapped

low-temperature superconductor because of recent STM results on the local density of states near magnetic impurities on niobium (Don Eigler's group, IBM Almaden). We explained the qualitative features of the data with a calculation based on a free-electron model (attached reprint). We have also explored many general features of the STM spectra around impurities on isotropically gapped superconductors using the remarkable energy resolution of the Koster-Slater technique (attached preprint). These include the spectra around non-magnetic and magnetic impurities in isotropically gapped superconductors with long and short coherence lengths.

A major theoretical conclusion of this work was that use of the correct detailed band structure of a material is critical for an accurate calculation of the STM spectra near an impurity. This is the case because the energy scale of impurities is atomic (eV), even though the energy of states created within the gap is of order the gap (meV). Hence the impurity potential mixes in states of the superconducting host which are very far away from the Fermi surface, and so a correct band-structure for those far away states is necessary. We are currently setting up a realistic band-structure model for niobium, based on thirty-four tight-binding parameters going out to the third nearest neighbors, which will allow us to compare quantitatively with the experimental data.

We have also produced an analytic model for the impurity within the *s*-wave superconductor based on a delta-function potential. Previous models assumed particle-hole symmetry, and thus produced pathological results for the local STM spectra. Our new analytic model has a single parameter which accounts for the actual case of no particle-hole symmetry. Thus the qualitative behavior of the local STM spectra can be understood without a detailed numerical calculation. This model is detailed in the attached preprint.

Once the utility of the new technique was demonstrated on isotropically

gapped superconductors we applied it to anisotropically gapped superconductors with a tight-binding band structure. The order parameters chosen were anisotropic *s*-wave and *d*-wave. STM spectra for magnetic and non-magnetic impurities in these superconductors have been calculated, and papers on these results are in preparation. For magnetic impurities we find that the sharp localized states within the gap which are present for isotropically gapped superconductors broaden into resonant states in the anisotropically gapped superconductors. These resonant states have highly asymmetric line-shapes and are present for both *d*-wave and anisotropic *s*-wave order parameters. Resonant states also are present for non-magnetic impurities for both types of order parameter.

The resonant states that we find should influence planar tunneling spectra when the impurity concentration is high enough. Since YBCO typically has a disordered surface, it is possible that these resonant states may be a cause of the two-gap features seen in many tunneling experiments. This type of structure in the density of states of the dirty superconductor should produce signatures in an assortment of macroscopic quantities, such as the specific heat and the magnetic penetration depth. In general, features which appear as double gaps may originate from an anisotropic order parameter (*s* or *d*) in the presence of impurities.

After submitting our results on anisotropically gapped superconductors we plan to investigate the other types of defects discussed in our proposal: step edges and vortices. An investigation of the step edge will be straightforward now that the single impurity problem is under control. Treating the twisting of the phase of the order parameter around a vortex core, however, will require a modification of the Koster-Slater technique. We expect this problem to be solvable without great difficulty.

LIST OF ATTACHED PAPERS

1. Michael E. Flatté and Jeff M. Byers, "Local Electronic Structure of a Single Magnetic Impurity in a Superconductor", *Physical Review Letters* **78**, 3761-3764 (1997). Development of the Koster-Slater technique and first application (magnetic impurities on niobium).
2. Michael E. Flatté and Jeff M. Byers, "Local Electronic Structure of Defects in Superconductors", *Physical Review B*, in press. Further development of the Koster-Slater technique and application to magnetic and non-magnetic impurities in isotropically gapped superconductors with long and short coherence lengths. Outline of technique for anisotropically gapped superconductors.

SEMINARS AND CONFERENCE TALKS

1. "Local Properties of Impurities in Superconductors", Solid State Seminar at Iowa State University, October 3, 1996.
2. "Local Properties of Impurities in Superconductors", Michael E. Flatté and Jeff M. Byers, American Physical Society March Meeting, Kansas City, March 21, 1997.
3. "Local Properties of Impurities in Superconductors", Solid State Seminar at Harvard University, April 24, 1997.

INVITED PAPERS

1. I have been invited to write a review article entitled "Local Properties of Defects in Superconductors" for the *Solid State Physics* series, published by Academic Press.

Local Electronic Structure of a Single Magnetic Impurity in a Superconductor

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We present the first three-dimensional, self-consistent calculation of the electronic structure near a strong classical magnetic impurity in a superconductor. Localized excited states are found within the energy gap which are half electron and half hole. The spatial structure of the positive-frequency (electronlike) spectral weight (or local density of states) can differ strongly from that of the negative frequency (holelike) spectral weight. The effect of the impurity on the continuum states above the energy gap is calculated with good spectral resolution for the first time. For sufficiently strong impurity potentials, the order parameter may change sign at the impurity site. [S0031-9007(97)03084-6]

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Magnetic impurities have a dramatic effect on superconductivity. Most works on the experimental and theoretical effects of magnetic impurities have focused on bulk thermodynamic quantities, such as the reduction of the superconducting transition temperature T_c with increasing magnetic impurity concentration [1]. Theoretical approaches to treat the thermodynamic effects of impurities include Born scattering (Abrikosov-Gor'kov theory) [2] and approximate solutions to all orders in the impurity potential for classical [3–5] and quantum [6,7] spins. A key issue addressed in Ref. [3] was the evolution of localized excited states (not obtainable in Born theory) around impurities into an impurity band, eventually leading to gapless superconductivity at high enough impurity concentrations. The effects of impurities on bulk properties have also been treated within a strong-coupling formalism (e.g., Ref. [8]), but only very recently self-consistently and beyond the Born approximation [7]. Concern about bulk properties in the above and related work did not extend to properties very near the impurity.

Among the first properties calculated in the vicinity of an impurity were the structures of screening clouds around a charged impurity [9,10] and a magnetic impurity [10,11] in a superconductor (characterized by exponentially decaying Friedel-like oscillations). The oscillation of the order parameter around a magnetic impurity was first evaluated without self-consistency [12,13]. A self-consistent calculation of the order parameter at the impurity and very far away for *weak* impurity potentials was done by Schlottmann [14]. Interest in local properties near impurities has been revived by advances in scanning tunneling microscopy (STM) near impurities embedded in a metallic medium [15].

The differential conductivity measured by a STM at a point \mathbf{x} , voltage V , and temperature T can be related to the local density of states (LDOS) at the tip location as follows:

$$\frac{dI(\mathbf{x}, V, T)}{dV} = \sum_{\sigma} \int_{-\infty}^{\infty} \frac{d\omega}{2\pi N_o} \frac{\partial n(\omega)}{\partial \omega} \text{Im} G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega). \quad (1)$$

Here σ is the spin direction of the electron ($\pm 1/2$), N_o is the normal-state density of states for each spin direction at the Fermi level, and dI/dV is in units of the normal-metal's differential conductivity. $n(\omega) = [1 + \exp(\beta(\omega - eV))]^{-1}$ where e is the electron's charge and $\beta = 1/T$. The LDOS is proportional to the imaginary part of the retarded Green's function fully dressed by the interaction of the electronic system with the impurity. The LDOS was calculated near impurities for superconductors with isotropic and anisotropic gaps within the Born approximation [16]. A self-consistent calculation of the LDOS of a vortex core in a superconductor was performed using the Schrödinger-like Bogoliubov-de Gennes (BdG) equations [17,18] and compared with measurements by an STM on superconducting NbSe_2 [19].

Recent STM results near a magnetic impurity [20] indicate that (1) discrete states are evident within the energy gap and the LDOS associated with them is asymmetric with voltage around $V = 0$, and (2) the LDOS becomes indistinguishable from the bulk density of states within a distance greater than the Fermi wavelength, but much less than the coherence length.

We will present a fully self-consistent calculation of the local electronic structure near a magnetic impurity, which is based on a Koster-Slater-like Green's function technique rather than the BdG equations [21]. Since the original application of the Green's function algorithm to localized vibrational modes [22], it has been applied to numerous problems, including conduction electrons in metals [23] and impurity states in magnets [24], but not to superconductors. The Green's function method works within a sphere whose radius is determined by the range of the inhomogeneous potential. In contrast, the BdG equations need to be solved in a large sphere whose radius is determined by the required spectral resolution. In essence we shall invert in real space the Dyson equation [25],

$$\begin{aligned} \int d\mathbf{x}'' [\delta(\mathbf{x} - \mathbf{x}'') - g_{\sigma}(\mathbf{x}, \mathbf{x}'', \omega) V(\mathbf{x}'')] G_{\sigma}(\mathbf{x}'', \mathbf{x}'; \omega) \\ = g_{\sigma}(\mathbf{x}, \mathbf{x}'; \omega), \end{aligned} \quad (2)$$

where

$$\mathbf{g}_\sigma(\mathbf{x}, \mathbf{x}'; \omega) = \begin{pmatrix} g_\sigma(\mathbf{x}, \mathbf{x}'; \omega) & f_\sigma(\mathbf{x}, \mathbf{x}'; \omega) \\ f_\sigma^*(\mathbf{x}, \mathbf{x}'; -\omega) & -g_{-\sigma}^*(\mathbf{x}, \mathbf{x}'; -\omega) \end{pmatrix}; \quad (3)$$

the same relationship exists among \mathbf{G}_σ , G_σ , and F_σ , and

$$\mathbf{V}(\mathbf{x}) = \begin{pmatrix} \sigma V_S(\mathbf{x}) + V_0(\mathbf{x}) & \delta\Delta(\mathbf{x}) \\ \delta\Delta(\mathbf{x}) & \sigma V_S(\mathbf{x}) - V_0(\mathbf{x}) \end{pmatrix}. \quad (4)$$

$\delta\Delta(\mathbf{x}) = \Delta(\mathbf{x}) - \Delta_o$, where $\Delta(\mathbf{x})$ is the position-dependent order parameter and Δ_o is the order parameter of the homogeneous superconductor. $V_S(\mathbf{x})$ is a localized, spin-dependent potential, such as one originating from a classical spin. $V_0(\mathbf{x})$ is a localized nonmagnetic potential. The quantization direction of the superconductor's spins is chosen parallel to the direction of the classical spin and so that spin-up electrons are attracted to the impurity. $G_\sigma \neq G_{-\sigma}$ due to the differences in the potential in Eq. (4). For a BCS superconductor with an isotropic order parameter in a parabolic band, for ω much smaller than the Fermi energy, the homogeneous retarded Green's functions are

$$\begin{aligned} g_\sigma(\mathbf{x}, \mathbf{x}'; \omega) &= -\frac{\pi N_o}{k_F r} e^{-\sqrt{\Delta_o^2 - \omega^2} r/\pi\xi\Delta_o} \\ &\quad \times \left(\cos k_F r + \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} \sin k_F r \right), \\ f_\sigma(\mathbf{x}, \mathbf{x}'; \omega) &= -\frac{\pi N_o \Delta_o}{k_F r \sqrt{\Delta_o^2 - \omega^2}} e^{-\sqrt{\Delta_o^2 - \omega^2} r/\pi\xi\Delta_o} \\ &\quad \times \sin k_F r, \end{aligned} \quad (5)$$

where $r = |\mathbf{x} - \mathbf{x}'|$, Δ_o is the order parameter, and $\xi = \hbar v_F / \pi \Delta_o$ is the coherence length. These expressions are valid for ω above and below Δ_o so long as the imaginary parts of both f_σ and g_σ are multiplied by $\text{sgn } \omega$. For a pairing strength γ , the self-consistency equation is

$$\delta\Delta(\mathbf{x}) = \gamma \sum_\sigma \int_{-\infty}^{\infty} d\omega n(\omega) \text{Im } F_\sigma(\mathbf{x}, \mathbf{x}; \omega) - \Delta_o. \quad (6)$$

One strength of this formalism is its reliance on the short-range nature of the inhomogeneous potential. The solution of Eq. (2) requires inverting the frequency-dependent real-space matrices $\mathbf{M}_\sigma(\omega) = \delta(\mathbf{x} - \mathbf{x}') - \mathbf{g}_\sigma(\mathbf{x}, \mathbf{x}'; \omega) \mathbf{V}(\mathbf{x}')$. For \mathbf{x}' 's where $\mathbf{V}(\mathbf{x}')$ is negligible, that portion of $\mathbf{M}_\sigma(\omega)$ is triangular and trivially invertible. Hence the numerical complexity of inverting $\mathbf{M}_\sigma(\omega)$ is determined by the range of $\mathbf{V}(\mathbf{x})$, which is governed by the range of the longest-range potential, which in this Letter will be $\delta\Delta(\mathbf{x})$.

As a special case of Eq. (2), the energies of the localized states within the gap correspond to those $\omega > 0$ where $\det \mathbf{M}_\sigma(\omega) = 0$. We model the impurity with a Gaussian $V_S(\mathbf{x})$ of range k_F^{-1} . Figure 1 shows the dependence of the localized state energies for the first two angular momentum channels on the strength of the magnetic potential, $v_s = \pi N_o |\sigma \int d\mathbf{x} V_S(\mathbf{x})|$. We note that a localized state implies local pair-breaking excitations of energy less than $2\Delta_o$. If

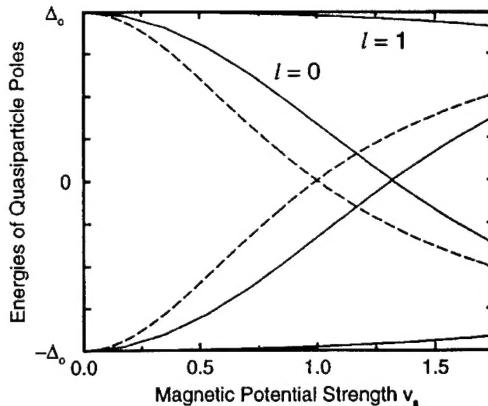


FIG. 1. Localized excited state poles for angular momentum channels $\ell = 0$ and 1 (solid lines). Dashed lines show results of Ref. [3]. At $v_s = v_s^*$, the character of the $\ell = 0$ state changes from spin up to spin down.

$\det \mathbf{M}_\sigma(\omega) = 0$, then $\det \mathbf{M}_{-\sigma}(-\omega) = 0$, so the poles of $G_\sigma(\omega)$ must come in symmetric pairs ($\sigma = \pm 1/2$) around $\omega = 0$. The quasiparticle state for small v_s corresponding to these poles consists of an electron ($\omega > 0$) in the spin-up (\uparrow) band (the band attracted to the classical spin) and a hole ($\omega < 0$) in the spin-down (\downarrow) band. A hole in the down band has spin up and is attracted to the classical spin. Spin is a good quantum number for the quasiparticle, which is spin up. The spatially integrated spectral weight is one-half each for the electron and hole components.

As the potential strength increases, the excitation energy of each angular momentum state ℓ decreases, and then at some critical value ($v_{s\ell}^*$) begins to increase. For $v_s > v_{s\ell}^*$ the *ground* state contains a spin-up quasiparticle bound to the classical spin [5]. The low-energy excitation now corresponds to removing that quasiparticle, which is a spin-down excitation. This qualitative behavior of the excitation energies can be extracted from the Shiba model [3], where the magnetic potential is modeled by a delta function at the impurity site. In Ref. [3] the zeros of $\mathbf{M}_\sigma(\omega)$ are found, neglecting the frequency symmetric component (whose significance is discussed below) of $\text{Re } g(\mathbf{x}, \mathbf{x}'; \omega)$. These poles are also shown in Fig. 1.

The decrease in the quasiparticle energy by the classical spin is proportional to the overlap of the atomic-scale potential and the impurity state. The state is, aside from an r^{-2} falloff, extended over a volume whose linear scale is the coherence length. Thus the overlap, and consequently the quasiparticle's energy reduction, is proportional to $v_s \Delta_o$. When this energy exceeds the energy required to create a quasiparticle (Δ_o), the ground state contains a quasiparticle. Hence v_s^* is approximately independent of Δ_o [3], and is determined primarily by the normal-state properties of the superconductor.

In the normal state the spectral weight of the spin-up band at the Fermi energy has a peak at the origin, while that of the spin-down band is pushed away from the origin. In the superconductor this asymmetry is evident

in the quasiparticle spectral weights. Figure 2 shows the calculated angular momentum $\ell = 0$ spectral weights in the up and down bands for two values of v_s , indicating that the asymmetry of the spatial structure of the spectral weights becomes more pronounced with increasing v_s . Also shown are the $\ell = 1$ spectral weights for $v_s = 0.875$. The asymmetric localized-state spectral weights produce differential conductivities which are asymmetric as well. Figure 3 shows differential conductivities for $v_s = 7/8$ as a function of the voltage and the distance from the impurity. By $r = 5k_F^{-1}$ the spectrum has recovered to the homogeneous (bulk) spectrum. The differential conductivity farther away can be easily recovered by constructing the self-consistent T matrix, $V(x')\delta(x' - x'') + V(x')G(x', x''; \omega)V(x'')$, directly from the dressed Green's functions.

The asymmetric behavior of the electronlike and holelike spectral weights due to the difference between the spin-up and spin-down band states does not emerge from the model of Ref. [3]. The Shiba model, due to the assumption of particle-hole symmetry implied by setting the symmetric real part of $g(\mathbf{0}, \mathbf{0}; \omega)$ to zero, predicts identical spatial dependences of the spectral weights for the up and down bands. As pointed out by Koster and Slater [23], proper treatment of the symmetric real part of $g(\mathbf{0}, \mathbf{0}; \omega)$ is essential for obtaining the local electronic properties around an impurity. The quantity $\text{Re } g(\mathbf{0}, \mathbf{0}; \omega)$ is simply the Kramers-Kronig transform of the density of states. The spatially dependent form of $\text{Re } g(x, x'; \omega)$ determines how the continuum states are distorted by the impurity potential to form the bound states. Neglecting the symmetric part of $\text{Re } g(x, x'; \omega)$ corresponds to a particle-hole symmetric density of states around the Fermi energy. The potentials considered here are locally strong, and mix in states of the homogeneous superconductor with energies much farther from the Fermi surface than Δ_o . Often a benign approximation when bulk properties are being calcu-

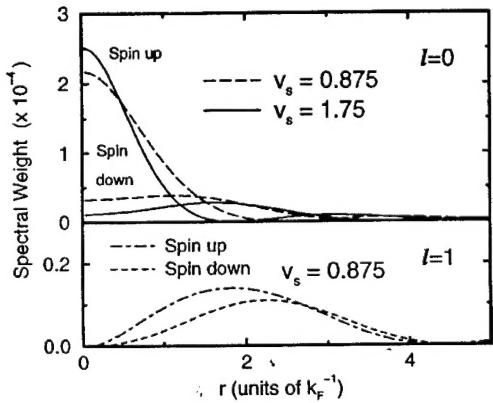


FIG. 2. Spectral weights for the $\ell = 0$ state's up-band component (electronlike for $v_s < v_s^*$) and down-band component (holelike for $v_s < v_s^*$) as a function of position for two values of the magnetic potential. Also shown are the spectral weights for the up and down bands for $\ell = 1$ for $v_s = 0.875$.

lated, particle-hole symmetry leads to pathological properties when calculating local electronic structure.

We introduce a new parameter into the Shiba model, α , which is the ω -symmetric part of $\text{Re}\langle g(\mathbf{0}, \mathbf{0}; \omega = 0) \rangle / \pi N_o$. The brackets indicate that $g(\mathbf{0}, \mathbf{x}; \omega = 0)$ has been averaged over a small volume given by the range of the potential modeled by the delta function. For $\alpha \neq 0$ the ratio for the localized quasiparticle of the spectral weight of the up band at the impurity, $A_{\uparrow}(\mathbf{0})$, to the spectral weight of the down band at the impurity, $A_{\downarrow}(\mathbf{0})$, is

$$\frac{A_{\uparrow}(\mathbf{0})}{A_{\downarrow}(\mathbf{0})} = \frac{1 + 2\alpha v_s + (1 + \alpha^2)v_s^2}{1 - 2\alpha v_s + (1 + \alpha^2)v_s^2}. \quad (7)$$

The introduction of α does not change the localized state energies qualitatively.

We now discuss the structure of $\Delta(\mathbf{x})$. Figure 4 shows $\delta\Delta(\mathbf{x})$'s for two values of v_s . While oscillating with wavelength $\sim \pi k_F^{-1}$, $\delta\Delta(\mathbf{x})$ falls off to a negligible potential within $10k_F^{-1}$. This justifies performing the numerical calculation in a sphere of that radius. A typical radial grid of 100 points provides a numerically robust solution. The self-consistent solution also depends on the value of $N_o \Delta_o / k_F^3$, which for niobium is 3.6×10^{-5} [26]. Relaxation of the order parameter on the atomic scale has also been seen in square-lattice calculations of the BdG equations for nonmagnetic impurities in quasi-two-dimensional d -wave superconductors [27], which should behave in a somewhat similar fashion to magnetic impurities in s -wave superconductors.

As shown in Fig. 4, for large values of v_s , $\Delta(\mathbf{x} = \mathbf{0}) < 0$. Sign changes in Δ , which could be seen in pair tunneling, have been suggested for magnetic impurities in

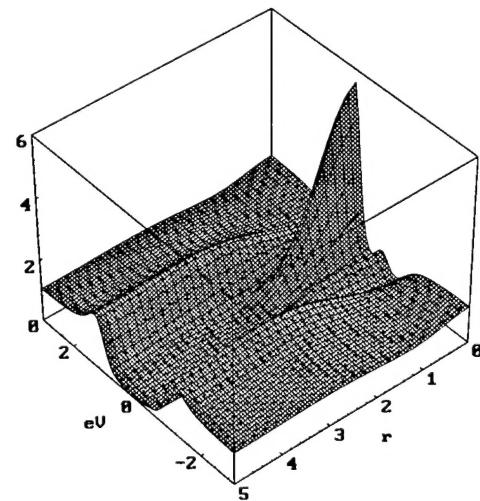


FIG. 3. Differential conductivity (dI/dV), relative to the normal metal as a function of the voltage (in units of Δ_o) and the distance from the impurity (in units of k_F^{-1}) for $v_s = 7/8$. The presence of the large peak near the origin on the positive-frequency side of the spectrum indicates that the impurity is not stronger than the critical strength ($v_s < v_{s0}^*$). $\beta = 7.5/\Delta_o$, which for niobium corresponds to $T \sim 2$ K.

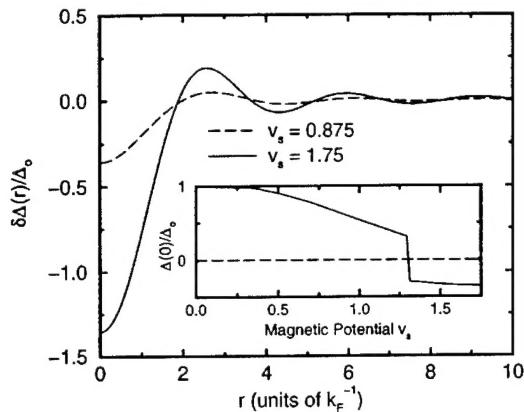


FIG. 4. Change in the local order parameter, $\delta\Delta(x)$, for two values of the magnetic potential. The change becomes negligible beyond $10k_F^{-1}$. Inset: Order parameter at the impurity, $\Delta(0)$, as a function of v_s , indicating a discontinuous change at v_{s0}^* .

the barriers of Josephson junctions [28]. Our sign change in $\Delta(0)$ occurs (at $T = 0$) precisely at v_{s0}^* . The symmetry of Eqs. (2)–(6) under $\omega \rightarrow -\omega$ and $s \rightarrow -s$ implies that $\text{Im } F_1(\mathbf{r}, \mathbf{r}, \omega) = -\text{Im } F_1(\mathbf{r}, \mathbf{r}, -\omega)$. As the pole in the spin-up band goes from electronlike ($\omega > 0$) to holelike ($\omega < 0$) and the pole in the spin-down band goes from holelike to electronlike, the contribution to $\Delta(0)$ from the localized state changes sign abruptly. $\Delta(0)$ as a function of v_s is shown in the inset of Fig. 4. At $T > 0$ the transition would be smoothed.

The behavior of $\Delta(0)$ as a function of v_s comes from the introduction at v_{s0}^* of a quasiparticle into the ground state of the system. The spin-up quasiparticle localized near the impurity in the ground state suppresses the local order parameter. Exciting the low-energy state (removing the spin-up quasiparticle) causes a pair to enter the condensate. We find that exciting the low-energy state for $v_s > v_{s0}^*$ increases $\Delta(0)$ and reduces the spin of the superconductor near the impurity. Typically the excitation of quasiparticles reduces $\Delta(x)$ (which is the case for $v_s < v_{s0}^*$).

We have calculated the electronic properties near a magnetic impurity embedded in a superconductor, and used these results to explain STM measurements around a magnetic atom on a surface. The asymmetry of the peak heights can arise from particle-hole asymmetry in the normal state of the superconductor. The recovery of the spectrum to its bulk structure within a few atomic spacings is due to the r^{-2} decay of the impurity state spectral weight, and the change in sign of the order parameter for sufficient v_s occurs because a quasiparticle is bound in the ground state. None of these aspects of the system change qualitatively at a surface. Thus the calculations presented here should have great value for the experimental (surface) configuration.

We wish to thank M. Salkola and A. V. Balatsky for pointing out Ref. [5] and clarifying Sakurai's picture of

one $\ell = 0$ localized state and the transition at v_s^* . During review we have become aware of related work by M. Salkola, A. V. Balatsky, and J. R. Schrieffer [29]. We wish to acknowledge conversations with B. A. Jones and A. Yazdani. M. E. F. wishes to acknowledge the Office of Naval Research's Grant No. N00014-96-1-1012. J. M. B. wishes to acknowledge a N.R.C. Postdoctoral Fellowship.

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Local Electronic Structure of Defects in Superconductors

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The electronic structure near defects (such as impurities) in superconductors is explored using a new, fully self-consistent technique. This technique exploits the short-range nature of the impurity potential and the induced change in the superconducting order parameter to calculate features in the electronic structure down to the atomic scale with unprecedented spectral resolution. Magnetic and non-magnetic static impurity potentials are considered, as well as local alterations in the pairing interaction. Extensions to strong-coupling superconductors and superconductors with anisotropic order parameters are formulated.

I. INTRODUCTION

Low-temperature superconductors almost always have a high concentration of non-magnetic impurities. Even in the dirty limit, however, where the mean free path is shorter than the coherence length, superconductivity endures¹. This phenomenon can be understood by generalizing BCS pairing² to be between degenerate Kramers partnered states in a time-reversal invariant system³. Magnetic impurities, which do break time-reversal invariance, have more profound effects on the superconductor in dilute concentrations than non-magnetic impurities, lowering the critical temperature T_c ⁴⁻⁶ and producing localized states within the gap⁷⁻¹³ which at sufficient concentrations hybridize to produce gaplessness¹⁴. In the course of the investigation of the effects of impurities on superconductivity during the last four decades, the primary emphasis has been the influence of impurities on bulk properties. These effects have been treated within a strong-coupling formalism (e.g. Ref. 15), but only very recently self-consistently and beyond the Born approximation¹⁶. The above work was primarily concerned with bulk or impurity-averaged characteristics and ignored the spatial structure of electronic properties near to the impurity.

Among the first local properties calculated in the vicinity of an impurity in a superconductor were the structures of screening clouds around a charged impurity^{17,18} and a magnetic impurity^{18,19} in a superconductor (characterized by exponentially-decaying Friedel-like²⁰ oscillations). The oscillation of the order parameter around a magnetic impurity was first evaluated without self-consistency²¹⁻²³. A self-consistent calculation of the order parameter at the impurity and very far away for weak impurity potentials was reported by Schlottmann²⁴ for magnetic impurities and Fetter²⁵ for non-magnetic impurities.

Interest in local properties near impurities in superconductors has been revived by the capability of scanning tunneling microscopy (STM) to perform localized

spectroscopic measurements. The spatially-dependent differential conductivity around a non-magnetic impurity at the surface of a superconductor has been considered theoretically²⁶ for both isotropic and anisotropic order parameters. This differential conductivity is proportional to the local density of states (LDOS) around the impurity. In Ref. 26 the impurity was modeled as a point defect, and spatial oscillations in the LDOS at various voltages were calculated. These oscillations are the superconductor's analog of oscillations in the LDOS created by step edges and defects on noble metal surfaces^{27,28}. The superconductor's LDOS oscillations would allow one to measure the anisotropy of the superconductor's order parameter. The conductance oscillations for voltages just above a gap minimum or maximum are strongly pronounced in the real-space directions corresponding to the momenta of the gap minimum and maximum. Calculations followed which considered sharp energy features in the scattering process, such as resonant states^{29,30}. Nevertheless, an important assumption of these calculations has remained unchecked in detail, that the electronic distortions induced by the impurity are local, including the deformation of the order parameter. Self-consistent calculations using the Bogoliubov-de Gennes (BdG) equations³¹ followed for two-dimensional systems³²⁻³⁵, but have been hampered by finite-size effects.

A magnetic impurity differs from a non-magnetic impurity in that a localized state exists around it^{7-13,36}. The first calculations of the LDOS of the localized and continuum states around a magnetic impurity were performed recently both with a simplified analytic model and numerically via the new self-consistent technique³⁷ described in this paper. Calculations of the LDOS of the localized state with angular momentum quantum number $\ell = 0$ were performed with a slightly different analytic model and numerically in two dimensions via the self-consistent BdG equations³⁸. These calculations were motivated by preliminary experimental results around a Gd atom on a niobium surface³⁹.

Another type of defect is a local change in the pairing interaction. The resulting spatially-dependent order parameter can then distort the LDOS. An order parameter suppression can even localize states, as in a vortex core^{40,41}.

The local electronic properties of all of these defects can be calculated self-consistently from the Gor'kov equation⁴² without further approximation with the new technique introduced in Ref. 37. Our technique for calculating the electronic structure around a defect in a superconductor is related to the Koster-Slater inversion techniques for determining the local electronic structure of impurities in metals^{43,44}. Since its original application to localized vibrational modes⁴⁵, this algorithm has been applied to numerous problems including deep levels in semiconductors⁴⁶ and impurity states in magnets⁴⁷. The new Koster-Slater technique separates space around the defect into two regions: the near field and the far field. The far field is a region distant enough from the defect that the potential is insignificant and the order parameter has relaxed back to its homogeneous value. The near field is the region close to the defect where the potential is finite or the order parameter is distorted. In essence, the Gor'kov equation that determines the Green's functions of the inhomogeneous superconductor is inverted in the real space region of the near field. This paper describes the technique in detail, expands on an analytic model introduced in Ref. 37 and reports several calculations of the properties of the defects described above.

Previous attempts to calculate the structure of the near field have often used the BdG equations³¹. Other theories of inhomogeneities, such as Ginzburg-Landau theory⁴⁸ or the Eilenberger equations⁴⁹, treat the spatial degrees of freedom as coarse-grained over the superconductor's coherence length. Coarse-grained approximations are not appropriate for considering electronic structure on the atomic scale near a defect. The BdG equations are generalized Schrödinger equations for the electron and hole wavefunctions of a quasiparticle, and are valid for a superconductor with an arbitrarily-varying order parameter, only constrained by the validity of BCS theory.

Unfortunately, these equations have significant practical difficulties as well. Despite qualitative success modeling STM measurements of a single vortex in superconducting NbSe₂⁵⁰, calculations of the electronic structure⁵¹⁻⁵⁵ using the BdG equations are hampered by the difference in energy scales between the Fermi energy and the order parameter. Since the BdG equations are solved numerically for a finite system, the difficulty of the calculation is determined by the necessary spectral resolution. The key energy scale which must be resolved is the superconducting gap. Thus, the numerical difficulty increases as the ratio of the Fermi energy to the gap becomes large. Hence the band structure assumed for the superconductor must be somewhat unrealistic (for Refs. 54, 55 the Fermi wavelength was approximately 100 Å, which is inappropriately large⁵⁶). This limitation extends to calculations of the interaction between a vor-

tex and an impurity⁵⁷, the characteristics of the vortex lattice^{58,59}, and work on a non-magnetic impurity³²⁻³⁵ and a magnetic impurity³⁸ in a two-dimensional *s*-wave or *d*-wave superconductor. In contrast, the computational requirements of the Koster-Slater technique are determined by the range of the impurity potential, rather than the necessary spectral precision.

In the Section II of this paper we first describe the BdG formalism for local defect potentials and then compare with the new, Koster-Slater formalism. Section III describes an analytic model, based on a delta-function potential, which reproduces some of the quantitative behavior of the numerical results. Section IV discusses the results of the numerical calculations for magnetic impurities, non-magnetic impurities, impurities incorporating both magnetic and non-magnetic potentials, and inhomogeneities in the pairing interaction. A heuristic picture of the electronic structure near these impurities will be presented here, and the calculations will be compared with the analytic model of Section III. Section V generalizes the formalism of Section II to the case of strong-coupling and anisotropically-paired superconductors.

II. FORMALISM

A. Bogoliubov-de Gennes Equations

To place our new formalism in context we will contrast it with the BdG equations, which are Schrödinger-like equations for the electron and hole components of the quasiparticle wavefunctions $u(\mathbf{x})$ and $v(\mathbf{x})$ respectively. These are, for a free electron band structure with mass m , the positive-energy (E) solutions to

$$\begin{aligned} \left[-\frac{(\hbar \nabla)^2}{2m} - E + V_0(\mathbf{x}) + \sigma V_S(\mathbf{x}) \right] u_\sigma(\mathbf{x}) + \Delta(\mathbf{x}) v_\sigma(\mathbf{x}) &= 0 \\ \left[\frac{(\hbar \nabla)^2}{2m} - E - V_0(\mathbf{x}) + \sigma V_S(\mathbf{x}) \right] v_\sigma(\mathbf{x}) + \Delta(\mathbf{x}) u_\sigma(\mathbf{x}) &= 0. \end{aligned} \quad (1)$$

Here σV_S is a position-dependent, spin-dependent potential, such as one originating from an impurity with a classical spin. V_0 is a position-dependent non-magnetic potential and $\Delta(\mathbf{x})$ is the inhomogeneous order parameter. $\Delta(\mathbf{x})$ can be chosen real since the defect potential is real⁶⁰. The quantization direction of the electronic spins in the superconductor ($\sigma = \pm 1/2$) is chosen parallel to the classical spin. A classical spin has no quantum dynamics, and cannot flip the quasiparticle spin. Hence spin is a good quantum number for the quasiparticles and only two coupled equations (Eqs. (1)) are required. The combinations $\sigma V_S \pm V_0$ have physical significance: $\sigma V_S + V_0$ is the potential felt by an electron of spin σ , while $\sigma V_S - V_0$ is the potential felt by a hole of spin σ .

The spatially-dependent order parameter is determined self-consistently:

$$\Delta(\mathbf{x}) = \sum_{n\sigma} \gamma(\mathbf{x}, E_{n\sigma}) u_{n\sigma}(\mathbf{x}) v_{n\sigma}^*(\mathbf{x}) \tanh\left(\frac{E_{n\sigma}}{2k_B T}\right) \quad (2)$$

where n labels the states for each spin σ , T is the temperature, and k_B is Boltzmann's constant. $\gamma(\mathbf{x}, E_{n\sigma})$ is the effective electron-electron interaction potential, which is

$$\begin{aligned} \gamma(\mathbf{x}, E_{n\sigma}) &= \gamma_0 & E_{n\sigma} < \hbar\omega_D \\ &= 0 & E_{n\sigma} > \hbar\omega_D. \end{aligned} \quad (3)$$

For a spherically-symmetric defect the wavefunctions are eigenstates of the angular momentum with quantum numbers ℓ and m . Typically the defect is placed in a sphere of radius R with appropriate boundary conditions. The value of R is determined by the spectral resolution necessary for accurately evaluating Eq. (2) and the spectral width of features measurable by (for example) the STM. The typical complications resulting from approximating an infinite system by a finite-size system apply, such as discrete states above the energy gap and the heavy investment of computer time required for large values of R . For example, in the calculations for the vortex in NbSe₂^{54,55}, $\epsilon_F/\Delta_o = 32$ was the largest ratio of the Fermi energy to the homogeneous order parameter considered. This value is unrealistic, and is a result of inappropriately fitting the coherence length and upper critical field of NbSe₂ to a free-electron model. A more realistic band structure⁵⁶ has a bandwidth to energy gap ratio at least an order of magnitude greater, and a highly anisotropic Fermi velocity. A system where the free-electron model is appropriate, such as niobium, has an $\epsilon_F/\Delta_o = 705$ and is numerically inaccessible for general potentials V_S and V_0 .

B. Koster-Slater Inversion Formalism

We now introduce a self-consistent method which works within a sphere whose radius is determined by the range of the defect's potential and utilizes the continuum spectrum of the homogeneous superconductor. In essence, we invert the Gor'kov equation in real space. The Gor'kov equation⁴² for a defect in a superconductor can be written in the Nambu formalism⁶¹ as:

$$\begin{aligned} \int d\mathbf{x}'' [\delta(\mathbf{x} - \mathbf{x}'') - \mathbf{g}(\mathbf{x}, \mathbf{x}''; \omega) \mathbf{V}(\mathbf{x}'')] \mathbf{G}(\mathbf{x}'', \mathbf{x}; \omega) \\ = \mathbf{g}(\mathbf{x}, \mathbf{x}'; \omega) \end{aligned} \quad (4)$$

where the inhomogeneous retarded Green's function,

$$\mathbf{G}(\mathbf{x}, \mathbf{x}'; \omega) = \begin{pmatrix} G_\uparrow(\mathbf{x}, \mathbf{x}'; \omega) & F(\mathbf{x}, \mathbf{x}'; \omega) \\ F^*(\mathbf{x}, \mathbf{x}'; \omega) & -G_\downarrow^*(\mathbf{x}, \mathbf{x}'; -\omega) \end{pmatrix}. \quad (5)$$

The elements of this matrix are

$$G_\uparrow(\mathbf{x}, \mathbf{x}'; \omega) = -i \int_{-\infty}^{\infty} dt e^{i\omega t} \theta(t) \langle 0 | \{ \psi_\uparrow(\mathbf{x}'; t), \psi_\uparrow^\dagger(\mathbf{x}; 0) \} | 0 \rangle, \quad (6)$$

$$\begin{aligned} F(\mathbf{x}, \mathbf{x}'; \omega) = \\ -i \int_{-\infty}^{\infty} dt e^{i\omega t} \theta(t) \langle 0 | \{ \psi_\uparrow(\mathbf{x}'; t), \psi_\downarrow(\mathbf{x}; 0) \} | 0 \rangle, \end{aligned} \quad (7)$$

$$\begin{aligned} F^*(\mathbf{x}, \mathbf{x}'; \omega) = \\ -i \int_{-\infty}^{\infty} dt e^{i\omega t} \theta(t) \langle 0 | \{ \psi_\downarrow^\dagger(\mathbf{x}'; t), \psi_\uparrow^\dagger(\mathbf{x}; 0) \} | 0 \rangle, \end{aligned} \quad (8)$$

$$\begin{aligned} -G_\downarrow^*(\mathbf{x}, \mathbf{x}'; -\omega) = \\ -i \int_{-\infty}^{\infty} dt e^{i\omega t} \theta(t) \langle 0 | \{ \psi_\downarrow^\dagger(\mathbf{x}'; t), \psi_\downarrow(\mathbf{x}; 0) \} | 0 \rangle. \end{aligned} \quad (9)$$

The explicit subscripts \uparrow and \downarrow do *not* refer to the spin of the excitation in the superconductor but rather to the spin band of the normal-state used to construct the excitation. The key concept is that the spin-*up* band contains both *up* electrons and *down* holes just as the spin-*down* band contains both *down* electrons and *up* holes. The convention employed here is standard in the theory of semiconductors where a spin-up electron excited above the Fermi energy leaves a spin-down hole below the Fermi energy. This is convenient for magnetic potentials since if spin-up electrons are attracted to a magnetic impurity spin-down holes should be repelled by the impurity. In the presence of a single classical impurity spin, the quasi-particle spin is a good quantum number despite electron-hole mixing. Our convention determines the composition of a spin-up quasiparticle to be part spin-up electron and part spin-up hole, rather than part spin-up electron and part spin-down hole.

For $\omega > 0$, $\mathbf{G}(\mathbf{x}, \mathbf{x}'; \omega)$ describes spin-up excitations, involving the mixing of electrons in the spin-up band with holes in the spin-down band. For $\omega < 0$, $\mathbf{G}(\mathbf{x}, \mathbf{x}'; \omega)$ describes spin-down excitations, involving the mixing of electrons in the spin-down band with holes in the spin-up band. Since spin is a good quantum number it is not necessary to use a 4×4 formalism, such as that of Ref. 62. The notation here has been simplified and improved relative to Ref. 37.

The homogeneous Green's function \mathbf{g} is independent of σ , so

$$\mathbf{g}(\mathbf{x}, \mathbf{x}'; \omega) = \begin{pmatrix} g(\mathbf{x}, \mathbf{x}'; \omega) & f(\mathbf{x}, \mathbf{x}'; \omega) \\ f^*(\mathbf{x}, \mathbf{x}'; \omega) & -g^*(\mathbf{x}, \mathbf{x}'; -\omega) \end{pmatrix}. \quad (10)$$

The potential

$$\mathbf{V}(\mathbf{x}'') = \begin{pmatrix} V_S(\mathbf{x}'') + V_0(\mathbf{x}'') & \delta\Delta(\mathbf{x}'') \\ \delta\Delta(\mathbf{x}'') & V_S(\mathbf{x}'') - V_0(\mathbf{x}'') \end{pmatrix} \quad (11)$$

where $\delta\Delta(\mathbf{x}) = \Delta(\mathbf{x}) - \Delta_o$ and V_S , V_0 , and $\Delta(\mathbf{x})$ have similar meaning above as in the BdG equations. The factor of one-half from the electron spin has been incorporated into the potential V_S . To be concrete, and without loss of generality, the spin direction attracted by the potential will be called spin up, and the spin direction repelled will be called spin down⁶³. The self-consistency equation for the order parameter is

$$\delta\Delta(\mathbf{x}) = \int_{-\infty}^{\infty} d\omega \gamma(\mathbf{x}, \omega) n(\omega) \text{Im}F(\mathbf{x}, \mathbf{x}; \omega) - \Delta_o \quad (12)$$

where $n(\omega)$ is the Fermi occupation function at temperature T .

Self-consistent spin-dependent and charge-dependent potentials can also be constructed for $\mathbf{V}(\mathbf{x})$ using the calculated spatial structure of the spin $s(\mathbf{x})$ and charge $\rho(\mathbf{x})$ around the defect,

$$s(\mathbf{x}) = \int_{-\infty}^{\infty} d\omega n(\omega) \left(-\frac{\text{Im}[G_{\uparrow}(\mathbf{x}, \mathbf{x}; \omega) - G_{\downarrow}(\mathbf{x}, \mathbf{x}; \omega)]}{\pi} \right) \quad (13)$$

$$\rho(\mathbf{x}) = \sum_{\sigma} \int_{-\infty}^{\infty} d\omega n(\omega) \left(-\frac{\text{Im}G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega)}{\pi} \right). \quad (14)$$

No calculations are reported for such potentials³¹ in this paper.

For a BCS superconductor with an isotropic gap in a parabolic band, for ω much smaller than the Fermi energy,

$$\begin{aligned} g(\mathbf{x}, \mathbf{x}'; \omega) &= -\frac{\pi N_o}{k_F r} e^{-\sqrt{\Delta_o^2 - \omega^2} r / \pi \Delta_o \xi} \left(\cos k_F r + \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} \sin k_F r \right) \\ f(\mathbf{x}, \mathbf{x}'; \omega) &= -\frac{\pi \Delta_o N_o}{k_F r \sqrt{\Delta_o^2 - \omega^2}} e^{-\sqrt{\Delta_o^2 - \omega^2} r / \pi \Delta_o \xi} \sin k_F r \end{aligned} \quad (15)$$

where $r = |\mathbf{x} - \mathbf{x}'|$ and N_o is the density of states for each spin at the Fermi level. The coherence length, $\xi = \hbar v_F / \pi \Delta_o$, where v_F is the Fermi velocity. These expressions are valid for ω above and below Δ_o so long as the imaginary parts of both f and g are multiplied by $\text{sgn}\omega$.

One strength of this formalism is its reliance on the short-range nature of the defect's potential. Solution of Eq. (4) requires inverting the frequency-dependent real-space matrix

$$\mathbf{M}(\mathbf{x}, \mathbf{x}'; \omega) = \delta(\mathbf{x} - \mathbf{x}') - \mathbf{g}(\mathbf{x}, \mathbf{x}'; \omega) \mathbf{V}(\mathbf{x}'). \quad (16)$$

The structure of $\mathbf{M}(\mathbf{x}, \mathbf{x}'; \omega)$ allows for a precise description of the difference between the near field and the far field. We require that the defect's potential $\mathbf{V}(\mathbf{x})$ is zero for $|\mathbf{x}| > R$. The space $|\mathbf{x}| > R$ belongs to the far field, whereas the space $|\mathbf{x}| \leq R$ belongs to the near field. We can then separate any real-space matrix \mathbf{A} symbolically into four pieces:

$$\mathbf{A} = \begin{pmatrix} \mathbf{A}^{n \rightarrow n} & \mathbf{A}^{n \rightarrow f} \\ \mathbf{A}^{f \rightarrow n} & \mathbf{A}^{f \rightarrow f} \end{pmatrix} \quad (17)$$

where n and f label the near-field region and far-field region, respectively. The particular example of \mathbf{M} is block-triangular:

$$\begin{aligned} \mathbf{M} &= \begin{pmatrix} \mathbf{I} - \mathbf{g}^{n \rightarrow n} \mathbf{V} & 0 \\ -\mathbf{g}^{f \rightarrow n} \mathbf{V} & \mathbf{I} \end{pmatrix} \\ \mathbf{M}^{-1} &= \begin{pmatrix} (\mathbf{I} - \mathbf{g}^{n \rightarrow n} \mathbf{V})^{-1} & 0 \\ \mathbf{g}^{f \rightarrow n} \mathbf{V} (\mathbf{I} - \mathbf{g}^{n \rightarrow n} \mathbf{V})^{-1} & \mathbf{I} \end{pmatrix}. \end{aligned} \quad (18)$$

It is clear from Eq. (18) that the computational effort in inverting \mathbf{M} , and thus finding the inhomogeneous electronic structure, is entirely determined by the complexity of inverting $\mathbf{M}^{n \rightarrow n}$. We also obtain the useful result that the electronic structure in the far field is easily determined once the electronic structure in the near field is known.

The local density of states in the near field is merely

$$\begin{aligned} -\frac{1}{\pi} \sum_{\sigma} \text{Im}G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega) &= -\frac{1}{\pi} \text{Im} \left\{ 2g(\mathbf{x}, \mathbf{x}; \omega) + \int d\mathbf{x}' \left[[(\mathbf{M}^{n \rightarrow n})^{-1}(\mathbf{x}, \mathbf{x}'; \omega)] \mathbf{g}(\mathbf{x}', \mathbf{x}; \omega) \right]_{11} \right. \\ &\quad \left. - \int d\mathbf{x}' \left[[(\mathbf{M}^{n \rightarrow n})^{(-1)*}(\mathbf{x}, \mathbf{x}'; -\omega)] \mathbf{g}^*(\mathbf{x}', \mathbf{x}; -\omega) \right]_{22} \right\} \end{aligned} \quad (19)$$

while in the far field it is

$$\begin{aligned} -\frac{1}{\pi} \sum_{\sigma} \text{Im}G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega) &= -\frac{1}{\pi} \text{Im} \left\{ 2g(\mathbf{x}, \mathbf{x}; \omega) \right. \\ &\quad + \int d\mathbf{x}' d\mathbf{x}'' \left[\mathbf{g}(\mathbf{x}, \mathbf{x}'; \omega) \mathbf{V}(\mathbf{x}') \left[(\mathbf{M}^{n \rightarrow n})^{-1}(\mathbf{x}', \mathbf{x}''; \omega) \right] \mathbf{g}(\mathbf{x}'', \mathbf{x}; \omega) \right]_{11} \\ &\quad \left. - \int d\mathbf{x}' d\mathbf{x}'' \left[\mathbf{g}^*(\mathbf{x}, \mathbf{x}'; -\omega) \mathbf{V}(\mathbf{x}') \left[(\mathbf{M}^{n \rightarrow n})^{(-1)*}(\mathbf{x}', \mathbf{x}''; -\omega) \right] \mathbf{g}^*(\mathbf{x}'', \mathbf{x}; -\omega) \right]_{22} \right\}. \end{aligned} \quad (20)$$

In the limit that the size of the near-field approaches a point, Eq. (20) is

$$-\frac{1}{\pi} \sum_{\sigma} \text{Im} G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega) = -\frac{1}{\pi} \text{Im} \left\{ 2g(\mathbf{x}, \mathbf{x}; \omega) + [\mathbf{g}(\mathbf{x}, \mathbf{0}; \omega) \mathbf{V}_{\text{eff}}(\omega) \mathbf{g}(0, \mathbf{x}; \omega)]_{11} - [\mathbf{g}^*(\mathbf{x}, \mathbf{0}; -\omega) \mathbf{V}_{\text{eff}}^*(-\omega) \mathbf{g}^*(0, \mathbf{x}; -\omega)]_{22} \right\}. \quad (21)$$

The inhomogeneous part of the right-hand side of Eq. (21) has the following form if the off-diagonal elements of $\mathbf{V}_{\text{eff}}(\omega)$ can be ignored (this is appropriate when self-consistency is not important):

$$-\frac{1}{\pi} \text{Im} \{ ([\mathbf{V}_{\text{eff}}(\omega)]_{11} - [\mathbf{V}_{\text{eff}}^*(-\omega)]_{22}) (g(\mathbf{x}, \mathbf{0}, \omega) g(\mathbf{0}, \mathbf{x}, \omega) - f(\mathbf{x}, \mathbf{0}, \omega) f(\mathbf{0}, \mathbf{x}, \omega)) \}. \quad (22)$$

Hence the spatially-dependent features reported in Ref. 26 are qualitatively retained in our more complete model. For a magnetic potential in the Born approximation there would be no such signal, since

$$\mathbf{V}_{\text{eff}}(\omega) = \begin{pmatrix} V_S & 0 \\ 0 & V_S \end{pmatrix} \quad (23)$$

so the distortions in the LDOS of the two spin directions are equal and opposite.

The local density of states under ideal conditions is directly measured by a scanning tunneling microscope. The differential conductivity measured at a point \mathbf{x} , voltage V and temperature T can be related to the local density of states at the tip location as follows:

$$\frac{dI(\mathbf{x}, V, T)}{dV} = \frac{e^2}{h} \int_{-\infty}^{\infty} d\omega \frac{\partial n_{\text{STM}}(\omega)}{\partial \omega} \sum_{\sigma} \left(\frac{\text{Im} G_{\sigma}(\mathbf{x}, \mathbf{x}; \omega)}{\pi} \right). \quad (24)$$

Here e is the charge of the electron and $n_{\text{STM}}(\omega)$ is the Fermi function in the STM tip

$$n_{\text{STM}}(\omega) = \left[1 + \exp \left(\frac{\omega - eV}{k_B T} \right) \right]^{-1}. \quad (25)$$

The local density of states is proportional to the imaginary part of the retarded Green's function fully dressed by the interaction of the electronic system with the impurity.

Certain features of the above equations simplify their numerical implementation. Each angular momentum channel constitutes an independent block-diagonal submatrix in $\mathbf{M}(\omega)$. Since the bare Green's functions in Eqs. (15) have analytic expansions in spherical harmonics, $\mathbf{M}(\omega)$ can be calculated quickly. These expansions are detailed in the Appendix.

The value of R is governed by the longest-range potential. In this paper that will be $\delta\Delta(\mathbf{x})$. We model V_S and V_0 with Gaussians of radius k_F^{-1} and evaluate Eq. (12) at $T = 0$. The $\delta\Delta(\mathbf{x})$ for $v_s = \pi N_o \int d\mathbf{x} V_S(\mathbf{x}) = 1.75$ and $v_0 = \pi N_o \int d\mathbf{x} V_0(\mathbf{x}) = 0$ is shown in Figure 1. While oscillating with wavelength $\sim \pi k_F^{-1}$, $\delta\Delta(\mathbf{x})$ falls off to a negligible potential within $10k_F^{-1}$. A typical radial grid

of 100 points provides a numerically robust solution. The self-consistent solution depends on the value of

$$N_o \Delta_o / k_F^3 = (2\pi^3 \xi k_F)^{-1} = (4\pi^2 \epsilon_F / \Delta_o)^{-1}, \quad (26)$$

which for niobium is 3.6×10^{-5} (Ref. 64). This is the single dimensionless parameter required to parametrize the free-electron model of a superconductor.

In contrast to the Koster-Slater technique described here, which exploits a physical distinction between the near field and the far field to accelerate the numerical calculation, the BdG equations treat the near-field and the far-field on the same level. A large R is desired for decent spectral resolution, but the possible size of R is limited by numerical constraints. Hence a numerical implementation of the BdG equations, by comparison, is typically slower and substantially less accurate than the Koster-Slater technique.

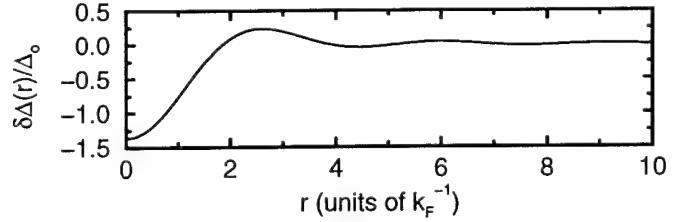


FIG. 1. Change in the local order parameter, $\delta\Delta(\mathbf{x})$, for a magnetic potential strength $v_s = N_o \int d\mathbf{r} V_S(\mathbf{r}) = 1.75$ as a function of the distance from the impurity r . The change becomes negligible beyond $10k_F^{-1}$.

III. ANALYTIC SOLUTION OF THE POINT POTENTIAL

A. Magnetic and Non-Magnetic Point Potentials

Approximating the local potential by a delta function

$$\mathbf{V}(\mathbf{x}) = \mathbf{V}\delta(\mathbf{x}) = \begin{pmatrix} V_S + V_0 & 0 \\ 0 & V_S - V_0 \end{pmatrix} \delta(\mathbf{x}) \quad (27)$$

leads to a simple expression for Eq. (4),

$$\begin{pmatrix} 1 - g(\mathbf{0}, \mathbf{0}; \omega)(V_S + V_0) & -f(\mathbf{0}, \mathbf{0}; \omega)(V_S - V_0) \\ -f^*(\mathbf{0}, \mathbf{0}; \omega)(V_S + V_0) & 1 + g^*(\mathbf{0}, \mathbf{0}; -\omega)(V_S - V_0) \end{pmatrix} \mathbf{G}(\mathbf{0}, \mathbf{0}; \omega) = \mathbf{M}^{n \rightarrow n}(\omega) \mathbf{G}(\mathbf{0}, \mathbf{0}; \omega) = \mathbf{g}(\mathbf{0}, \mathbf{0}; \omega). \quad (28)$$

In principle $\mathbf{M}(\omega)$ can be found from the Green's functions in Eq. (15), however there is a divergence in the real part of $g(r; \omega)$ as $r \rightarrow 0$. This divergence is coped with in Ref. 8 by discarding the divergent piece. This approximation is essentially an assumption of *strict* particle-hole symmetry (not merely linearizing $\epsilon(k)$ around ϵ_F). We now discuss the effects of this approximation on the local properties of the system in the *normal* state. The heuristics will be simpler for the normal state, but the conclusions also apply to the superconducting state.

B. Particle-Hole Symmetry in the Normal State

In order to focus on the spin-dependent potential, the non-magnetic potential will be set to zero. The normal state properties can be obtained from Eq. (15) for $\Delta_o = 0$. That yields a Green's function appropriate for an outgoing wave:

$$g(\mathbf{x}, \mathbf{x}'; \omega) = -\frac{\pi N_o}{k_F r} e^{ik_F r}, \quad (29)$$

where ω is considered close to the Fermi surface so the change in momentum due to $\omega \neq 0$ is negligible. The *inhomogeneous* local density of states for a delta-function potential using this Green's function is unphysical, since it diverges at $r = 0$. Now we examine the local density of states when the divergent real part of g is ignored.

$$-\frac{1}{\pi} \text{Im}G_\sigma(\mathbf{x}, \mathbf{x}; \omega) = N_o \left(1 + \left[\frac{(\pi N_o V_S)^2}{1 - (\pi N_o V_S)^2} \right] \frac{\sin^2 k_F r}{(k_F r)^2} \right). \quad (30)$$

This expression yields the pathological behavior that the local density of states near a spin-dependent potential is exactly the same for spin-up electrons as for spin-down electrons. A spatial response in the LDOS to a spin-dependent potential that is identical for up and down spin electrons *only* occurs for certain band structures with ϵ_F at special energies (e.g. half-filled tight-binding models). The local density of states for $v_s = \pi N_o |V_S| = 0.1$ is plotted in Figure 2 as a function of distance from the potential. Although it is somewhat distorted from its homogeneous value, it does not show the spin-dependent asymmetry of a more realistic potential. An exact calculation for a Gaussian of range k_F^{-1} is also shown in Figure 2 for comparison.

A more realistic approach to coping with the divergence in Eq. (29) without yielding the pathological result of Eq. (30) is to average the real part of g over a range given by the assumed range of the potential. That yields a finite value for the local density of states at the potential, but does not control the behavior for small r .

To perform that task, we consider a "muffin-tin" Green's function. This function has the form

$$\begin{aligned} g(\mathbf{x}, \mathbf{x}'; \omega) &= -i \frac{\pi N_o}{k_F r} e^{ik_F r} & r > R_o \\ &= -i\pi N_o - \pi\alpha & r < R_o \end{aligned} \quad (31)$$

where α is the average of the real part of g over the range of the potential, and R_o is chosen so that the spatially-integrated spectral weight of the Green's function is unchanged (as required by probability conservation). We show in Figure 2 the local density of states calculated with this Green's function. In particular, the asymmetry in the response of spin-up electrons and spin-down electrons to a spin-dependent delta-function potential is governed by this phenomenological parameter α . The agreement with the exact solution is good at the origin and far from the impurity. The muffin-tin Green's function is discontinuous, unfortunately, but yields a better approximation of the response of the system than the particle-hole symmetric approximation (discarding the real part of the Green's function).

It is also possible to generate an asymmetry between spin-up and spin-down electrons by adding a non-

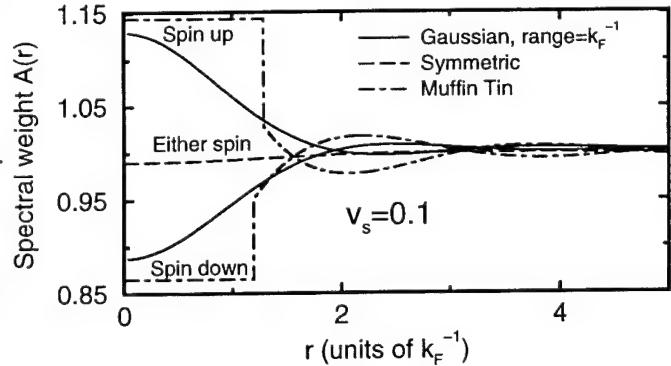


FIG. 2. Spectral weight in the normal state around a magnetic impurity with $v_s = 0.1$ as a function of the distance from the impurity r . The solid lines are exact solutions for spin-up (attracted by the impurity) and spin-down (repelled by the impurity) electrons for a Gaussian potential of range k_F^{-1} in a metal with a free-electron dispersion relation. The dashed line, which is the same for both spin-up and spin-down electrons, is the calculated spectral weight for the particle-hole symmetric delta-function potential model. The dot-dashed line is the result for a delta-function potential calculated using the muffin-tin Green's functions. The muffin-tin Green's functions fix the pathological result of the symmetric model that the spectral weight is the same for spin-up and spin-down electrons. The muffin-tin parameter, $\alpha = 0.704$, is determined by the range of the Gaussian potential and is therefore not a free parameter.

magnetic potential v_0 with the v_s to parametrize the impurity, but still maintaining a particle-hole symmetric band structure. Since the response of a particle-hole symmetric system to a potential does not depend on the sign of that potential, the v_0 is required to distinguish between electrons and holes. An electron feels a potential $v_s + v_0$, whereas a hole feels a potential $v_s - v_0$. This additional non-magnetic potential only breaks particle-hole symmetry locally (within the range of the potential), whereas for a realistic band structure the particle-hole symmetry is broken everywhere in the solid.

For a Gaussian potential with range a ,

$$\alpha = \frac{2}{\sqrt{\pi} k_F a} \left(1 + \sum_{n=2}^{\infty} \frac{1}{(2n-3)!!} \left(-\frac{(k_F a)^2}{2} \right)^{n-1} \right). \quad (32)$$

For the Gaussian potentials numerically calculated in this paper, $a = k_F^{-1}$, so $\alpha = 0.704$.

C. Self-consistency within the Analytic Model

As seen from Figure 1, the distortion of the order parameter is short-ranged around an impurity. We may then consider the effect of the order-parameter distortion on the electronic structure to be parametrized by an effective delta-function potential $\delta\Delta\delta(\mathbf{x})$ similarly motivated to the delta-function potentials for the magnetic and non-magnetic potentials. The potential $\mathbf{V}(\mathbf{x})$ is changed in the following way:

$$\mathbf{V}(\mathbf{x}) = \mathbf{V}\delta(\mathbf{x}) = \begin{pmatrix} V_S + V_0 & \delta\Delta \\ \delta\Delta & V_S - V_0 \end{pmatrix} \delta(\mathbf{x}). \quad (33)$$

The relative effect of the $\delta\Delta$ compared to the other two potentials is likely to be small for the potentials considered in this paper. Typically $N_o V_S / k_F^3 \sim 1$ or $N_o V_0 / k_F^3 \sim 1$, and for niobium $N_o \Delta_o / k_F^3 = 3.6 \times 10^{-5}$. Even for a small coherence length of $\xi = 10k_F^{-1}$, $N_o \Delta_o / k_F^3 = 1.6 \times 10^{-3}$. For convenience we define $\delta_0 = \pi N_o \delta\Delta$.

D. Energies of localized states in the Superconductor

The energies of the localized states of angular momentum ℓ correspond to the positive energies $\omega_\ell = |\Omega|$, where

$$\det \mathbf{M}^{n \rightarrow n}(\Omega) = 0, \quad (34)$$

and the solution is traced to the ℓ -channel block of \mathbf{M} (see Appendix). For the analytic model, $\mathbf{M}^{n \rightarrow n}(\omega)$ is the matrix shown in Eq. (28), where $g(\mathbf{0}, \mathbf{0}; \omega)$ in the superconducting state is constructed similarly to that of the normal state,

$$g(\mathbf{0}, \mathbf{0}; \omega) = -\pi N_o \left(\alpha + \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} \right). \quad (35)$$

The anomalous Green's function is given by Eq. (15), since it does not have a divergence problem as $r \rightarrow 0$,

$$f(\mathbf{0}, \mathbf{0}; \omega) = -\pi N_o \left(\frac{\Delta_o}{\sqrt{\Delta_o^2 - \omega^2}} \right). \quad (36)$$

This analytic model only has localized states in the $\ell = 0$ angular momentum channel, as expected for a delta-function potential. Those energies are

$$\omega_o = \left| \frac{v_s \delta_0 \pm [(2v_s \delta_0)^2 - 4(v_s^2 + \gamma^2)(\delta_0 - \gamma^2)]^{1/2}}{v_s^2 + \gamma^2} \right| \Delta_o \quad (37)$$

where

$$\gamma = [(1 + \alpha^2)(v_s^2 - v_0^2 - \delta_0^2) - 2\alpha v_0 - 1] / 2. \quad (38)$$

Eq. (37) reduces to a result obtained by Shiba⁸ when $v_0 = \alpha = \delta_0 = 0$, a result obtained by Rusinov⁹ when $\alpha = \delta_0 = 0$, a result obtained by Salkola, Balatsky and Schrieffer³⁸ when $\alpha = 0$, and a result obtained by us³⁷ when $v_0 = \delta_0 = 0$.

One solution of Eq. (37) is a spin-up quasiparticle and the other is a spin-down quasiparticle. There may be only one real solution to Eq. (37); then only one $\ell = 0$ localized state exists around the impurity. This occurs for large v_s . When $v_s = 0$ the localized states are due to order parameter suppression, and the energies of the two spin states are degenerate. This follows from time-reversal symmetry in the absence of a magnetic potential. For small v_s the two energies are split by an amount

$$\Delta\omega_o = \frac{2v_s \delta_o \Delta_o}{v_s^2 + \gamma^2}. \quad (39)$$

E. Spectral Weight Asymmetry in the Analytic Model

A spin-up quasiparticle consists of amplitudes for a spin-up electron (electron in a spin-up state), and a spin-up hole (electron missing from a spin-down state). Therefore the spectral weight of a spin-up localized state will be divided between an electron-like pole in the spin-up band at $\omega = \omega_o$ (with weight $A_\uparrow(\mathbf{r}; \omega_o)$) and a hole-like pole in the spin-down band at $\omega = -\omega_o$ (with weight $A_\downarrow(\mathbf{r}; -\omega_o)$). These two types of excitation are independently resolvable by a scanning tunneling microscope since at positive sample voltage relative to the tip, the STM places electrons in the sample, whereas at negative

sample voltage the STM places holes in the sample. We define the energy of the pole in the spin-up band to be ω_\uparrow and in the spin-down band to be ω_\downarrow . Even though ω_o is always positive, ω_\uparrow can be positive or negative, and $\omega_\uparrow = -\omega_\downarrow$.

The spatial structure of the spectral weights of the spin-up band and spin-down band components of the localized state are given by

$$\begin{aligned}
A_\sigma(\mathbf{r}; \omega) &= \frac{\pi N_o \Delta_o}{2v_s} \delta(\omega - \omega_\sigma) \frac{\sqrt{\Delta_o^2 - \omega^2}}{\Delta_o} \left[\frac{(v_0 - v_s)\Delta_o^2 + (v_0 + v_s)\omega^2 + (v_0 + v_s)\alpha^2(\Delta_o^2 - \omega^2)}{\Delta_o^2} \right. \\
&\quad \left. + \frac{2(v_0 + v_s)\alpha \omega \sqrt{\Delta_o^2 - \omega^2} - (1 + \alpha^2)(v_0^2 - v_s^2) \left(\alpha(\Delta_o^2 - \omega^2) + \omega \sqrt{\Delta_o^2 - \omega^2} \right)}{\Delta_o^2} \right] \quad r < R_o \\
&= \frac{\pi N_o \Delta_o}{(k_F r)^2} \frac{(\Delta_o^2 - \omega^2)^{\frac{3}{2}}}{2v_s \Delta_o^3} e^{-\frac{2r}{\pi \xi} \left(\frac{\sqrt{\Delta_o^2 - \omega^2}}{\Delta_o} \right)} \delta(\omega - \omega_\sigma) \times \\
&\quad \left[\left(v_s \frac{\Delta_o^2 + \omega^2}{\Delta_o^2 - \omega^2} - v_0 + (v_s^2 - v_0^2) \left\{ \alpha - \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} \right\} \right) \sin^2(k_F r) \right. \\
&\quad + (v_s + v_0) \left(1 + (v_s - v_0) \left\{ \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} - \alpha \right\} \right) \cos^2(k_F r) \\
&\quad \left. + 2(v_s + v_0) \left(v_0 - v_s + [1 - \alpha(v_0 - v_s)] \frac{\omega}{\sqrt{\Delta_o^2 - \omega^2}} \right) \sin(k_F r) \cos(k_F r) \right] \quad r > R_o \tag{40}
\end{aligned}$$

IV. COMPARISON WITH NUMERICAL RESULTS AND DISCUSSION

The above expression is set up for use as a muffin-tin Green's function. The construction of such a Green's function requires that the spectral weight integrates to one. However, if $R_o \rightarrow 0$, the above expression can be integrated over all space, and still yields one-half for the spin-up band and one-half for the spin-down band to order Δ_o/ϵ_F . Hence the quasiparticle is one-half electron and one-half hole. This provides additional confidence in the above expressions.

The frequency-integrated weight at the defect of the two types of excitation can be calculated within the analytic model, and is (for $\delta_0 = 0$)

$$\begin{aligned}
A_\uparrow(\mathbf{r} = \mathbf{0}) &= \frac{1 + 2\alpha(v_0 - v_s) + (1 + \alpha^2)(v_0 - v_s)^2}{1 + 2\alpha(v_0 + v_s) + (1 + \alpha^2)(v_0 + v_s)^2} \tag{41} \\
A_\downarrow(\mathbf{r} = \mathbf{0}) &=
\end{aligned}$$

For a spin-down quasiparticle there is an electron-like pole in the spin-down band and a hole-like pole in the spin-up band and the relative weight is still given by the expression above. This expression for $v_0 = 0$ was reported in Ref. 37.

In the following section these results are compared with the numerical calculations of properties in the superconducting state and in the normal state.

A. Non-Magnetic Impurity

Even strong non-magnetic impurities at moderate concentrations will not suppress the critical temperature of a superconductor³. Nevertheless, it was recognized early on²⁵ that the local order parameter may be affected. In Ref. 25 the effect of the non-magnetic impurity was calculated in the far field by modeling the impurity potential with a phase shift. The phase shifts were evaluated for two models: a spherical square-well potential and a delta-shell potential. Self-consistency was ignored by only focusing on regions far from the impurity where the change in the order parameter is small compared to its homogeneous value. The order parameter change due to the impurity was found to oscillate with the Fermi wavelength, and decay as r^{-2} .

Figure 3 shows the spectral weight $A(\mathbf{r}; \omega)$ at several frequencies above the energy gap near a strong non-magnetic impurity with a Gaussian potential of range k_F^{-1} and strength $v_0 = 7/8$, calculated self-consistently for a superconductor with $\xi k_F = 449$ (niobium). The spectral weights are suppressed to approximately 30% of their homogeneous value at the center of the potential. Only continuum states are shown since no localized states

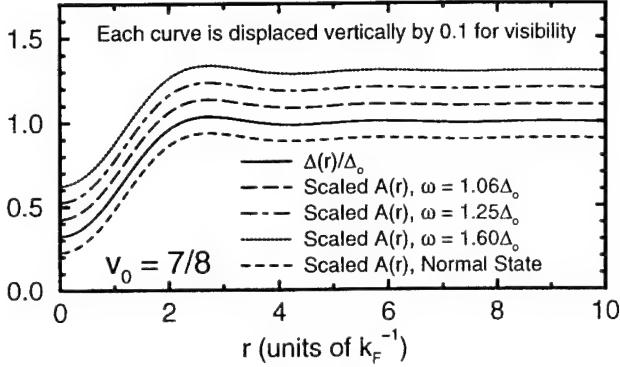


FIG. 3. Comparison of the spectral weight in the superconducting state at several frequencies to the spectral weight in the normal state and the spatial dependence of the order parameter. All of these curves are indistinguishable, but they have been displaced for better visibility. The spectral weights are normalized to their value at large distances from the impurity r , and the order parameter is normalized to the homogeneous order parameter Δ_o . This figure indicates that normal-state properties drive the spatially-dependent features of the superconductor's spectrum.

were found for this potential. The curves showing the spectral weight have been displaced from each other so that they may be distinguished. Also shown displaced in Fig. 3 is the spectral weight in the normal state, normalized to the spectral weight in the homogeneous metal. All the quantities plotted in Fig. 3 are *identical* to the accuracy of the calculation. Fig. 3 is an illustration of a relationship between the spectral weight in the normal state and the spectral weight in the superconducting state,

$$A(\mathbf{r}; \omega) = \frac{A_n(\mathbf{r})}{2N_o} A_{sc}(\omega) \quad (42)$$

where $A_n(\mathbf{r})$ is the spectral weight in the inhomogeneous *normal* state for energies near the Fermi surface and $A_{sc}(\omega)$ is the *homogeneous* superconductor's spectral weight as a function of frequency. $2N_o$ is the normal state's spectral weight far from the impurity. This expression is valid for small r and small ω , the regime of interest for STM on a superconductor. For ω of order Δ_o , Eq. (42) is valid for $r < \xi$.

We further illustrate the relationship of Eq. (42) in Figure 4(a), which shows the LDOS for this non-magnetic impurity as a function of voltage and position calculated from Eq. (24) with $T = \frac{2}{15}\Delta_o/k_B$. This temperature corresponds to 2K for niobium. There is no change in the energy *gap* due to this non-magnetic potential. Figure 4(a) shows that it is merely the local amplitude of the spectral weight which is reduced — this would manifest itself in a locally reduced oscillator strength for an optical transition, or the reduction in the tunneling current for an STM, which is directly proportional to the LDOS

shown in Figure 4(a). Figure 4(b) is an identical calculation for a shorter coherence length, $\xi = 10k_F^{-1}$. There appears to be little difference, although here a localized state exists near the continuum.

Figures 3 and 4(ab) show for the non-magnetic impurity that the normal-state electronic structure determines the *spatial* dependence of the superconductor's $A(\mathbf{r}; \omega)$ for all frequencies including near the energy gap. The potential strength of the impurity is orders of magnitude greater than Δ_o , and thus locally mixes in states far

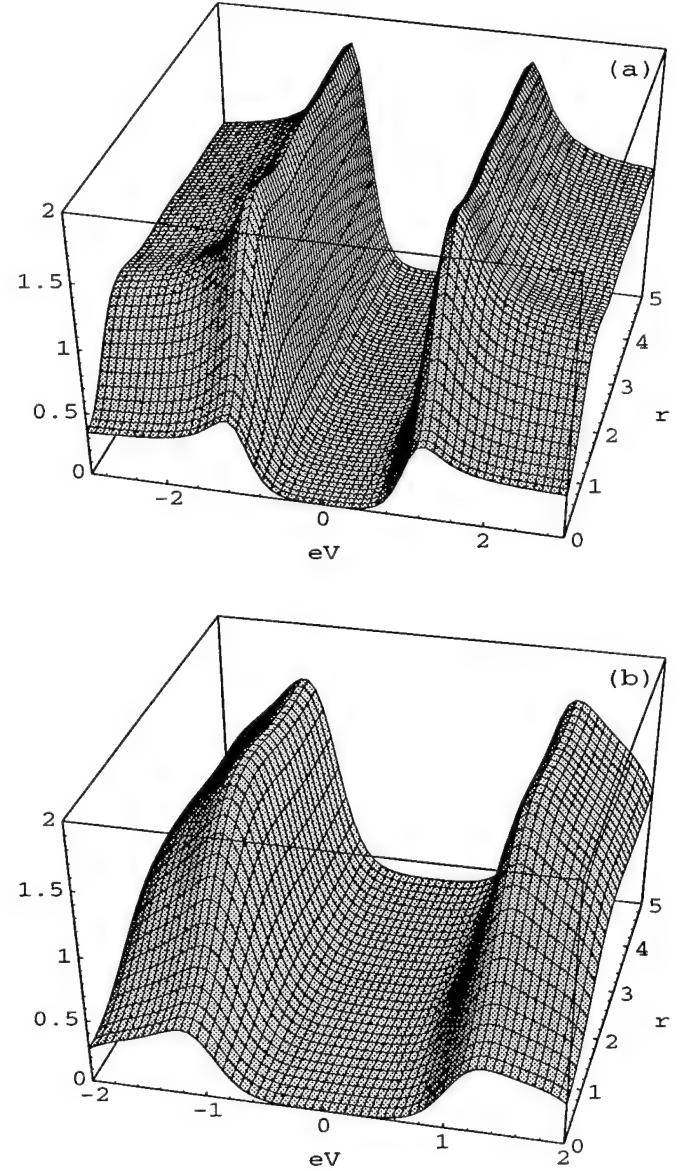


FIG. 4. Differential conductance (LDOS) as a function of voltage and position calculated around a non-magnetic impurity with $v_0 = 0.875$. The spectrum is suppressed substantially in the vicinity of the impurity. The temperature is $\Delta_o/7.5k_B$, which for niobium corresponds to about 2K. (a) A coherence length appropriate for niobium, $\xi k_F = 449$. (b) A much shorter coherence length, $\xi k_F = 10$.

from the Fermi surface in the homogeneous metal. These states are required to construct probability densities which are suppressed by 70% near the impurity. The spatial structure of the spectral weight in the normal state is essentially identical to that seen in Fig. 3 over an energy range around the Fermi surface which is orders of magnitude greater than Δ_o .

Once the normal-state band structure has been distorted by the presence of the non-magnetic impurity, superconductivity is a small perturbation within a narrower frequency range. The formation of the gap in the single-particle excitation spectrum in the superconducting state is characterized by the “mixing” of electron and hole amplitudes to form quasiparticles near the gap edge. These quasiparticles, therefore, are constructed from single-particle eigenstates of the metal which have already been strongly distorted by the impurity potential.

Equation (42) has important implications for spectroscopy on a superconductor, for one of the procedures for normalizing spectra taken at different lateral positions on a superconducting surface is to assume that the LDOS at a particular voltage much larger than Δ_o is the same. This is an attempt to correct for possible changes in the tip-surface distance upon moving the tip laterally. A small change in the tip-surface distance can have a strong effect on the tunneling current. Unfortunately this procedure will prevent an experiment from seeing changes in the LDOS due to a non-magnetic impurity, including the conductance oscillations described in Ref. 26.

We now discuss the properties of the order parameter. $\Delta(\mathbf{r})$ is self-consistently determined, and is shown in Fig. 3 for small r to be identical in spatial structure to the normal-state spectral weight,

$$\frac{\Delta(\mathbf{r})}{\Delta_o} = \frac{A_n(\mathbf{r})}{2N_o} = \frac{A(\mathbf{r}; \omega)}{A_{sc}(\omega)}. \quad (43)$$

Since a non-magnetic potential repulsive to electrons attracts holes, and $\Delta(\mathbf{r})$ depends equally on electron and hole amplitudes, one might expect a non-magnetic potential to have little effect on the spatial dependence of the order parameter. However the allowable maximum spectral density of holes depends on the spectral density of the electron band where the holes reside, so if most electrons are excluded from the site, holes will be effectively excluded as well. To emphasize this point we note that the scaled anomalous spectral weight $\text{Im}F(\mathbf{r}, \mathbf{r}; \omega)$ is identical to the scaled $A(\mathbf{r}; \omega)$ for the frequencies shown in Fig. 3 (and for all relevant frequencies for the self-consistency equation Eq. (12)). Since $\text{Im}F(\mathbf{r}, \mathbf{r}; \omega)$ is proportional to the product of electron and hole amplitudes, and $A(\mathbf{r}; \omega)$ is proportional to the electron amplitude squared, the spatial structure of the electron and hole spectral weights must be similar. They are, since the normal-state spectral weight $A_n(\mathbf{r}; \omega)$ is roughly frequency independent around the Fermi energy over an energy range much greater than Δ_o .

We now comment on the lack of localized states near the non-magnetic impurity for $\xi k_F = 449$ and the small binding energy for the quasiparticle for $\xi k_F = 10$. The suppression of the order parameter near the impurity may be considered to form an attractive off-diagonal potential which may bind quasiparticles. Localized states created by order-parameter suppression would be doubly degenerate, due to the two possible spin states (see Eq.(37)). Since the quasiparticle is half hole and half electron, if the electron part is attracted and the hole part is repelled, one might expect the effects of such a non-magnetic potential on the quasiparticle to cancel. However the binding energy of the localized state is an order of magnitude smaller ($\omega_o = (1 - 2 \times 10^{-4})\Delta_o$) than that found in Section IV.D for a suppressed order parameter via pairing suppression. This may be explained by the well-known repulsive effect (quantum reflection) of a strong attractive potential on a quantum-mechanical particle. We find also that in the case of the magnetic impurity that a large enough non-magnetic potential of either sign will suppress the binding of a quasiparticle to the impurity. We note here that the ratio of the non-magnetic potential to the off-diagonal potential ($\delta\Delta(\mathbf{r})$) is much larger ($v_0/N_o\Delta_o \sim 10^4$) than the ratios of the non-magnetic potentials to the magnetic potentials considered below.

B. Magnetic Impurity

Recently, we have presented calculations of the LDOS (and thus the differential conductivity in an STM experiment) in the vicinity of a magnetic impurity³⁷. These calculations indicated that the spatial structure of the electron amplitude of the localized state differed strongly from the hole amplitude of the localized state. A further result was that the spectrum should recover to the homogeneous spectrum within a few atomic spacings. Similarly motivated calculations of the LDOS due to the $\ell = 0$ localized state have been presented since then³⁸, although these calculations did not address the continuum LDOS. The two models used in Ref. 38 were (1) a δ -function model solved using particle-hole symmetric Green’s functions, but not self-consistently, and (2) self-consistent calculations for a two-dimensional tight-binding s -wave superconductor within the BdG equations. The first method can only model the normal-state properties properly for a particle-hole symmetric band structure, such as at the Van Hove singularity in a two-dimensional tight-binding band structure. The second method must contend with numerical finite-size effects, which make it difficult to calculate the continuum states. A result obtained from the first method which is only true for special band structures is that the spatial structure of the electron and hole components of the quasiparticle are the same. The authors of Ref. 38 did raise the possibility of an additional non-magnetic potential as a source

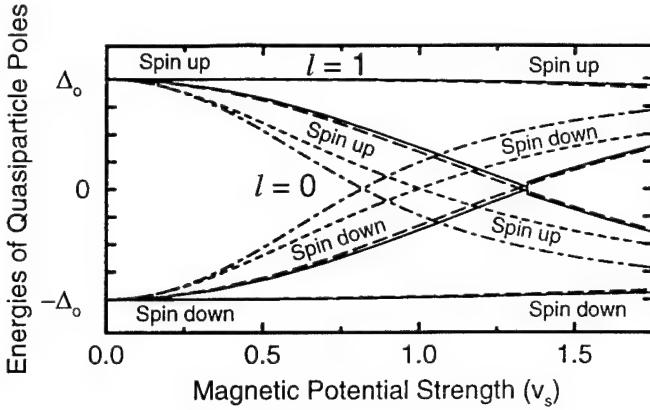


FIG. 5. Energies of the quasiparticle poles as a function of the magnetic potential strength v_s for the first two angular momentum states around the impurity. The spin-up and spin-down labels refer to the band that the excitation resides in — an excitation with negative energy (hole-like) in a spin-up band is a spin-down hole. The solid line corresponds to $\xi k_F = 10$, the long-dashed line to $\xi k_F = 449$, the short dashed line to the symmetric model of Shiba, and the dot-dashed line to the result calculated with muffin-tin Green's functions and $\alpha = 0.704$. At a critical value of $v_s = v_{s0}^*$, the up poles cross to negative energies and the down poles cross to positive energies, indicating a change in the character of the ground state. The kink evident in the solid and long-dashed lines is real, and due to the discontinuous (at $T = 0$) change in $\Delta(x)$ at v_{s0}^* . The unimportance of self-consistency can be gauged by the small difference between the short-coherence length result and the long-coherence length result.

of electron-hole amplitude asymmetry in the spatial structure of the localized state. We found³⁷ and will explore below that there is, for realistic band structures, electron-hole asymmetry without a non-magnetic potential.

We will begin with a discussion of the energies of the localized states around a magnetic impurity and the spin character of those states. Solutions to Eq. (34) can be evaluated numerically. Figure 5 shows the dependence of the energies of the localized state poles for the first two angular momentum channels on the strength of the magnetic potential. Results for a short coherence length ($\xi = 10k_F^{-1}$) are shown (solid line) as well as results for a long coherence length ($\xi = 449k_F^{-1}$). The localized quasiparticle state for small v_s is the spin state attracted to the classical spin, which we will label up (\uparrow)⁶³. As the potential strength increases, the excitation energy of each angular momentum state decreases. At some critical value $v_{s\ell}^*$ the localized state becomes a spin-down excitation, the energy changes abruptly, and then increases with increasing v_s . This behavior can be extracted from the analytic model (Eq. (37)) as well.

Also shown in Figure 5 are the analytic results for the pole energies for $\alpha = 0$ (Ref. 8) and $\alpha = 0.704$. The

muffin-tin model is no better than the particle-hole symmetric model in predicting the localized state energies. The muffin-tin model will prove more successful at predicting the asymmetry between the electron and hole amplitudes of the localized state.

The unimportance of self-consistency for determining the energies of the localized states can be gauged by the small difference between the short-coherence length result and the long-coherence length result. The most important feature it determines in Fig. 5 is the size of the discontinuity in the localized state energy at v_{s0}^* . This discontinuity is due to a discontinuous change in the order parameter at this magnetic potential strength, a result pointed out in Ref. 37 which will be discussed more below.

A magnetic impurity affects each member of a pair oppositely. An up-spin electron is attracted to the impurity and a down-spin electron is repelled. The difference in energy between the two time-reversed electron states leads to a locally suppressed pairing. Therefore, the energy needed to break a pair and create a localized quasiparticle with angular momentum ℓ in the vicinity of the magnetic impurity, when $v_s < v_{s\ell}^*$, is reduced from $2\Delta_0$ to $\Delta_0 + \omega_\ell$. One member of the broken pair is a delocalized spin-down quasiparticle at the gap edge (with energy Δ_0). The other member of the broken pair is a *localized* spin-up quasiparticle with energy ω_ℓ . For $v_s > v_{s\ell}^*$ there is a spin-up quasiparticle with angular momentum ℓ in the ground state of the superconductor⁶⁵. An essential point about the ground state of a superconductor containing classical magnetic impurities is that at $T = 0$ when $v_s < v_{s\ell}^*$ (for all ℓ) the ground state is composed entirely of paired electrons. When $v_s > v_{s\ell}^*$ for any ℓ then the $T = 0$ ground state contains localized quasiparticles as well as pairs and there are new low-energy excitations including the re-formation of a pair as well as the excitation of a localized quasiparticle to a higher energy localized or continuum state.

Evident from Eq. (5) is that the electron pole involves entirely single-particle states within the spin-up band for $v_s < v_{s\ell}^*$, but involves entirely states within the spin-down band for $v_s > v_{s\ell}^*$. The hole-like pole always involves single-particle states in the opposite spin band from the electron-like pole. The source of the quasiparticle amplitude for the various poles is indicated in Fig. 5.

Figures 6(a), 6(b), and 6(c) show LDOS results for $v_s = 0.52$, 0.875 , and 1.75 respectively. They show the state split off from the continuum, with a larger electron-like amplitude than hole-like amplitude (6(a)), and then lower in energy with an increased electron/hole asymmetry (6(b)). Finally the larger peak becomes hole-like (6(c)). The asymmetry between the electron-like and hole-like peaks becomes more pronounced as v_s increases. We note that the larger peak is always associated with the spin-up band, whereas the other is associated with the spin-down band. Despite the apparent differences in peak size, the spatially integrated electron spectral weight of the quasiparticle is equal to the spatially inte-

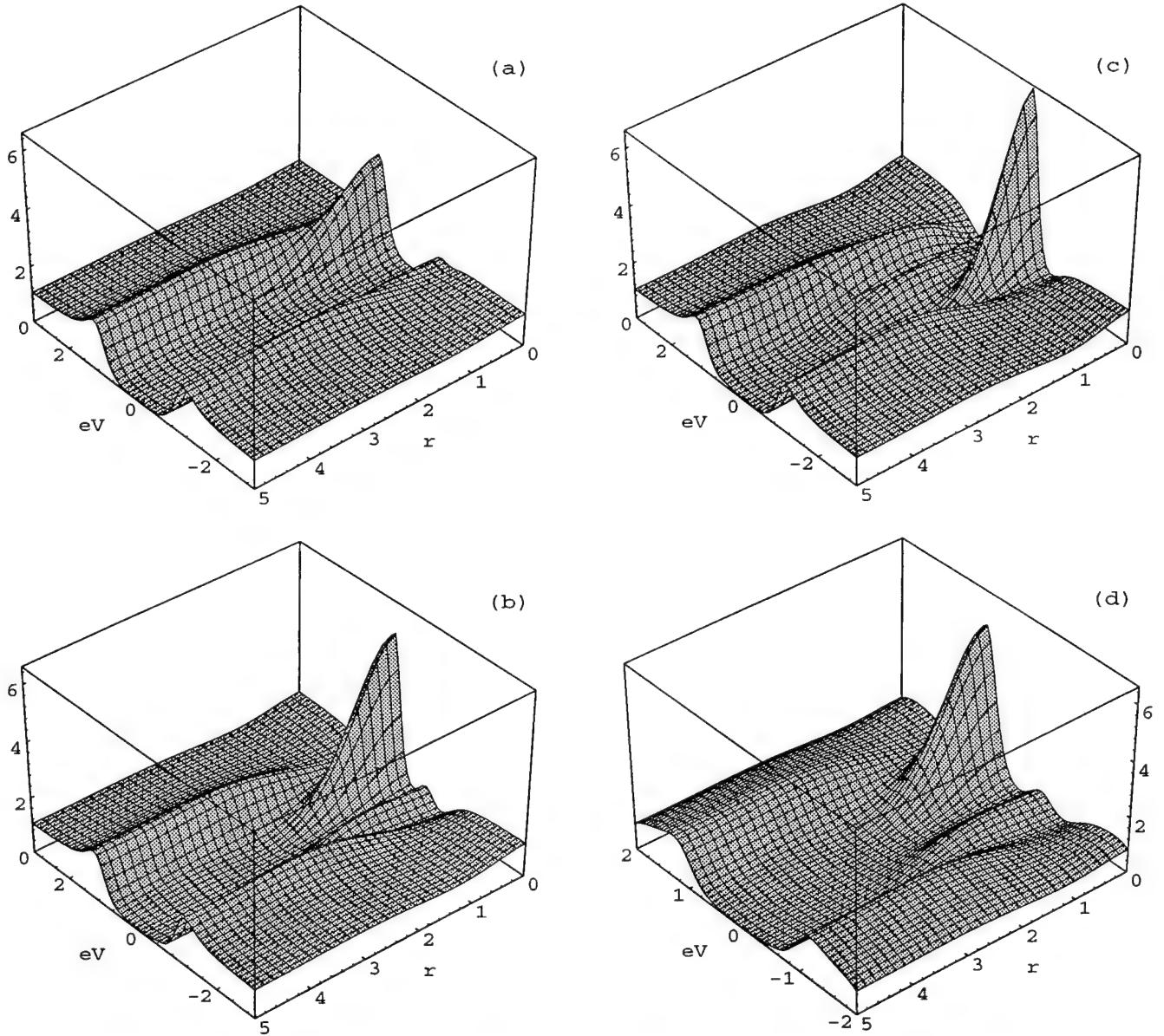


FIG. 6. Differential conductance (LDOS) calculated around a magnetic impurity with (a) $v_s = 0.5$, $\xi k_F = 449$, (b) $v_s = 0.875$, $\xi k_F = 449$, (c) $v_s = 1.75$, $\xi k_F = 449$, (d) $v_s = 0.875$, $\xi k_F = 10$. All are calculated with $k_B T = \Delta_o / 7.5$. In the progression from (a) to (c) the asymmetry between the two peaks increases and the higher peak moves to lower energies, eventually becoming hole-like.

grated hole spectral weight. The localized quasiparticle is always half electron and half hole for all potentials examined here. For $v_s < v_{s0}^*$ the spin-up band amplitude is electron-like and the spin-down band amplitude is hole-like. At v_{s0}^* (1.32 for niobium), the spin-up component becomes hole-like and the spin-down component becomes electron-like, as required by the change in the spin of the excitation.

Figure 6(d) shows the LDOS for a markedly different coherence length, $\xi = 10k_F^{-1}$, and $v_s = 0.875$. It is evaluated for the same values of $\Delta_o/k_B T$ as Figs. 6(abc) and looks almost identical to Fig. 6(b). Since the localized state is broadened by temperature through Eq. (24), this

is a manifestation of the proportionality of the spectral weight to $N_o \Delta_o$ (Eq. (40)). Figure 7 shows the spectral weight at the origin for the $\ell = 0$ state and for the $\ell = 1$ state at its first maximum for $v_s = 0.875$ as a function of the inverse of the coherence length, which is proportional to $N_o \Delta_o$. Figure 8 shows the spectral weight for $\xi = 449k_F^{-1}$ as a function of v_s for the spin-up and spin-down poles at the origin for the $\ell = 0$ state and at the first maximum for the $\ell = 1$ state. It is clear that a non-magnetic potential is not necessary to obtain an electron-hole asymmetry.

In Figure 9 we show the asymmetry at the impurity as a function of v_s for two values of ξ — a long coherence

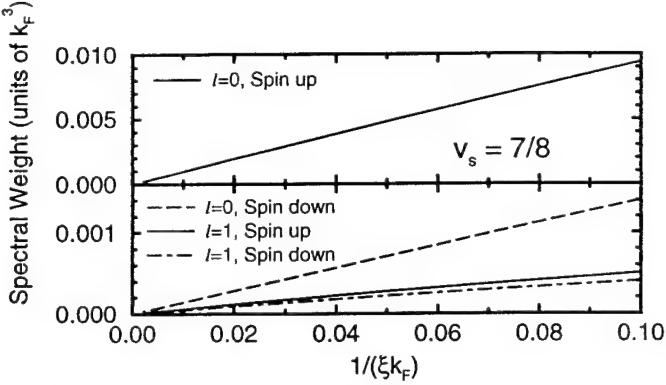


FIG. 7. Spectral weight at the impurity ($r = 0$) for the first angular momentum channel, $\ell = 0$, and at the first maximum for the second angular momentum channel, $\ell = 1$, for poles in both the spin-up and spin-down bands, as a function of the inverse coherence length, showing a linear behavior. The magnetic potential strength is $v_s = 0.875$.

length appropriate for niobium, and a short coherence length. From Figure 7 it should be evident that the asymmetry is not sensitive to ξ . It is, however, predicted extremely well by the normal-state spin-up and spin-down band spectral weight asymmetry at the impurity (also shown in Figure 9). We can therefore conclude that as with the non-magnetic impurity (Figure 4), the spatial structure of the spectral weight is a normal-state property. We further show in Figure 10 for $v_s = 0.875$ the $\ell = 0$, spin-up band and spin-down band projections of the normal-state spectral weight to compare with the localized state spin-up band and spin-down band spectral weights for two values of ξk_F . The normal-state and long-coherence length calculation are practically indistinguishable. The insets show $r^2 A(r)$, which removes the rapid power-law decay of the state. The localized states for all angular momenta ℓ will decay as the power law r^{-2} . For the short-coherence length calculation the

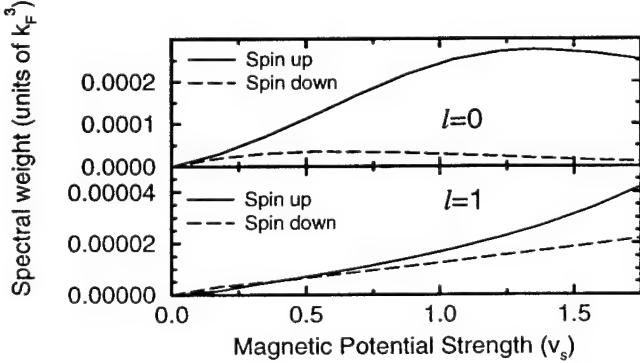


FIG. 8. Spectral weight at the impurity ($r = 0$) for the $\ell = 0$ channel for $\xi k_F = 449$ as a function of magnetic potential strength for poles in both the spin-up and spin-down bands. The spectral weight in the spin-up band pole of the $\ell = 0$ localized state saturates at large v_s . Also shown are the spectral weights at the first maximum for the $\ell = 1$ localized states.

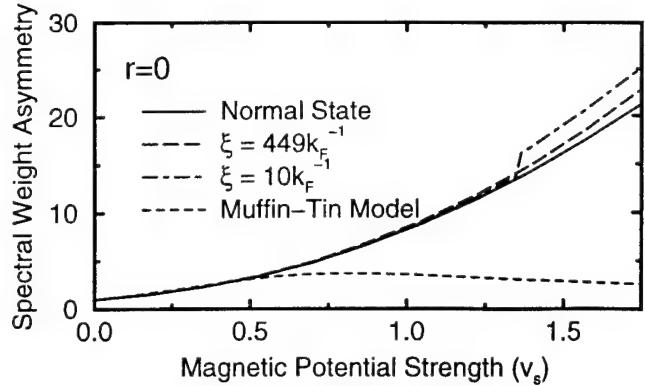


FIG. 9. Ratio of the spectral weight in the spin-up band and in the spin-down band at the impurity ($r = 0$) as a function of magnetic potential strength. This is plotted for the normal state $\ell = 0$ projected Green's functions (solid line) as well as for the localized states for niobium ($\xi k_F = 449$, long dashed line), for $\xi k_F = 10$ (dot-dashed line), and for the muffin-tin model. The muffin-tin model is only successful for $v_s < 0.5$, but that is due to a breakdown in describing the normal state. The normal-state electronic structure is a good predictor of the superconductor's electronic structure for the entire range of v_s .

effect of an exponential envelope is also visible. In the analytic result the exponential envelope should have a range $R = \pi \xi / 2 \sqrt{1 - (\omega_o / \Delta_o)^2}$, which corrects to better than 1% the discrepancy in Figure 10. The power-law

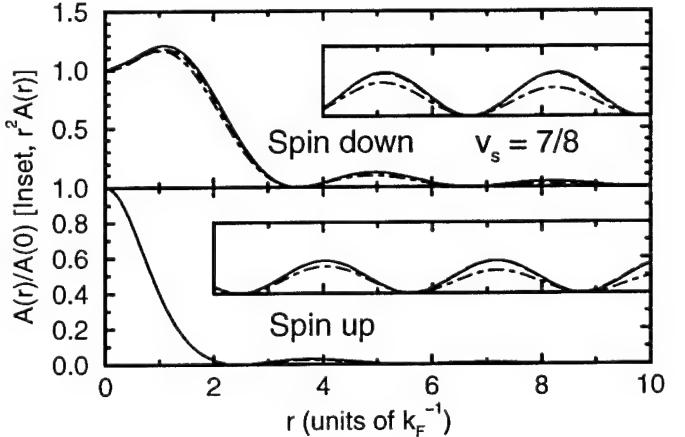


FIG. 10. Spectral weights for $\ell = 0$ localized state in the up and down bands for $v_s = 0.875$. The solid line is the normal state $\ell = 0$ projected spectral weight, the long-dashed line is the localized state in a superconductor with $\xi k_F = 449$, and the dot-dashed line is for $\xi k_F = 10$. The inset shows the spectral weight multiplied by r^2 to remove the algebraic decay. The normal-state and long-coherence length results are practically indistinguishable. The deviation shown in the short-coherence length superconductor's spectral weight is fit to within 1% by the exponential decay factor described in the text. Hence the spatial structure of the spectral weight of the superconductor's localized state is well-predicted by the normal state spectral weight.

fall-off and exponential envelope can be seen directly from Eq. (15) and Eq. (40).

We can summarize these comments with a general equation, similar in concept to that for the non-magnetic impurity, Eq. (42). That is, for a localized quasiparticle state of spin σ' , the spectral weight of a localized state with angular momentum ℓ would be

$$A_\sigma(\mathbf{r}; \omega_\ell) = BA_{n\sigma}(\mathbf{r}, \ell)e^{-(\frac{2r}{\pi\xi})\sqrt{1-(\frac{\omega_\ell}{\Delta_0})^2}}\delta(\omega - \sigma\sigma'\omega_\ell), \quad (44)$$

where B is a normalization factor so that the spectral weight of the state integrates to one, and $A_{n\sigma}(\mathbf{r}, \ell)$ is the angular-momentum ℓ projection of the *normal-state* spectral weight in the spin σ band. We note that for small r there is an approximate relationship between the superconducting state's spectral weight and the normal state's spectral weight in each spin band,

$$\frac{1}{2E} \int_{-E}^E d\omega A_\sigma(\mathbf{r}; \omega) = A_{n\sigma}(\mathbf{r}), \quad (45)$$

where $\Delta_0 \ll E \ll \epsilon_F$. This, in connection with Eq. (44), implies a dependence on the normal-state structure of the continuum spectral weight around the magnetic impurity.

We now return to the structure of $\Delta(\mathbf{x})$. This quantity, which is not directly observable, formed the focus of several investigations of the local structure around a

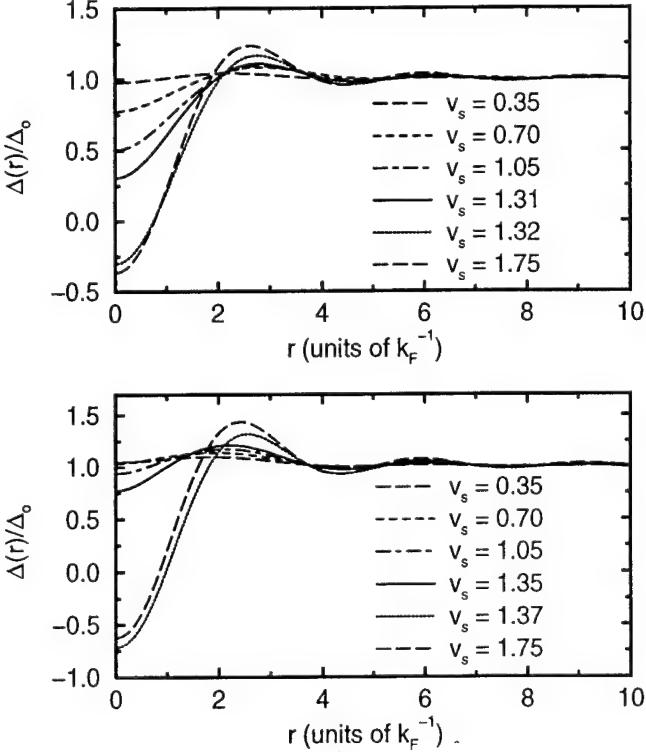


FIG. 11. Order parameters as a function of distance from the impurity r calculated for several magnetic potential strengths for (a) $\xi k_F = 449$ and (b) $\xi k_F = 10$. In both cases there is a discontinuous change in the order parameter when v_s passes through the critical strength v_{s0}^* .

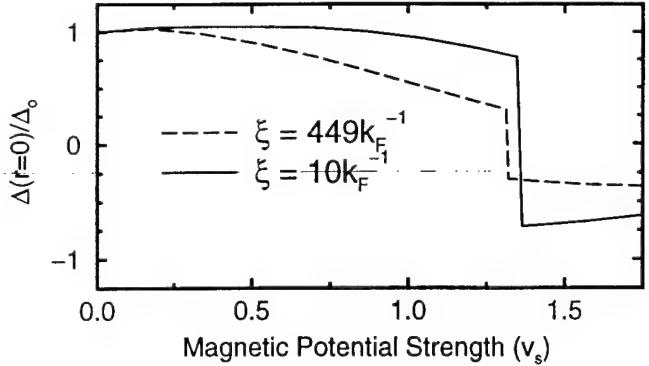


FIG. 12. Order parameters at the impurity ($r = 0$) as a function of magnetic potential strength v_s for two values of the coherence length. The discontinuity in the order parameter at v_{s0}^* is much larger for the short-coherence-length superconductor.

magnetic impurity. The oscillation of the order parameter around a magnetic impurity was first evaluated without self-consistency²¹⁻²³. A self-consistent calculation of the order parameter at the impurity and very far away for *weak* impurity potentials was done by Schlottmann²⁴.

As shown in Figure 1, for large values of v_s , $\Delta(\mathbf{x} = 0) < 0$. Sign changes in Δ , as seen in pair tunneling, have been suggested for magnetic impurities in the barriers of Josephson junctions⁶⁶⁻⁶⁸. The sign change in $\Delta(\mathbf{0})$ occurs (at $T = 0$) precisely at v_{s0}^* . Due to the spin and frequency symmetries of Eqs. (4-11), the anomalous spectral weight $\text{Im}F(\mathbf{r}, \mathbf{r}, \omega)$ associated with the spin-up pole is always equal and opposite to the anomalous spectral weight associated with the spin-down pole. As the pole in the spin-up band goes from electron-like ($\omega > 0$) to hole-like ($\omega < 0$) and the pole in the spin-down band goes from hole-like to electron-like the contribution to $\Delta(\mathbf{0})$ changes sign abruptly at $T = 0$. The $\Delta(\mathbf{r})$ resulting from several values of v_s and two values of the coherence length are shown in Figure 11(ab). The discontinuity at v_{s0}^* is more pronounced for shorter coherence lengths since the localized state's spectral weight is more concentrated at the impurity (Eq. (44)). $\Delta(\mathbf{0})$ as a function of v_s is shown in Figure 12 for two values of the coherence length. At $T > 0$ the transition would be smoothed somewhat.

The behavior of $\Delta(\mathbf{0})$ as a function of v_s comes from the introduction at v_{s0}^* of a quasiparticle into the ground state of the system. The spin-up quasiparticle localized near the impurity in the ground state suppresses the local order parameter. For time-reversal invariant potentials one cannot make $\Delta(\mathbf{r})$ negative by inserting a single quasiparticle, since the suppression from one quasiparticle is cancelled by the lack of suppression from its unexcited Kramers doublet partner. For a spin-dependent potential, however, the anomalous spectral weight near the impurity may be almost entirely contributed by the single low-energy localized state. When a quasiparticle is present in the ground state, the ground state has spin $\frac{1}{2}$

up^{65,38} and a negative $\Delta(\mathbf{r})$ ³⁷. Exciting the low-energy state for $v_s > v_{s0}^*$ removes the spin-up quasiparticle, and therefore *increases* $\Delta(0)$, whereas excitation of quasiparticles typically reduces $\Delta(\mathbf{x})$ (which is the case for $v_s < v_{s0}^*$). Also, exciting the low-energy state *reduces* the induced spin of the superconductor at the impurity.

C. Combined Magnetic and Non-Magnetic Potentials

We now discuss the addition of a non-magnetic potential to the magnetic potential. It had been suggested³⁸ that introducing a v_0 with a v_s would provide electron-hole asymmetry. We find that it does change the asymmetry, which we show in Figure 13 for a particular v_s , but that once again this is a normal-state property. The relationship between the normal-state spectral weights and the superconducting-state spectral weights of Eq. (44) still holds. Introducing v_0 also alters the localized-state energies (see Eq. (37)), which we show in Figure 14 for $v_s = 0.875$ and $\xi k_F = 449$. The presence of a non-magnetic potential may affect the value of v_{s0}^* ³⁸. We show in Figure 15 a partial diagram of the ground state as a function of the parameters v_s and v_0 for $\xi k_F = 449$. We note that the boundary between the two ground states is not shifted much from $\xi k_F = 449$ to $\xi k_F = 10$, hence the condensation energy is not very significant in determining this boundary. This fact, and the observation that the energy of the localized state (in the absence of self-consistency) relative to the homogeneous order parameter does not depend on any parameter of the superconducting state (Eq. (37)) suggests that an understanding of the transition at v_{s0}^* may be assisted by normal-state concepts. We hope to address this issue in a future

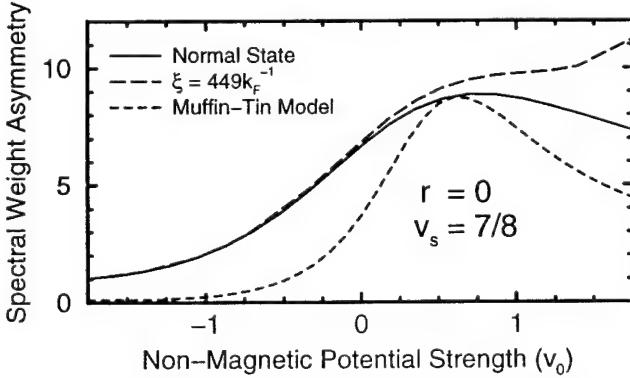


FIG. 13. Ratio of the spectral weight at the impurity in the spin-up band to the spin-down band for $v_s = 0.875$ and $\xi k_F = 449$ as the non-magnetic potential v_0 varies. In a similar result to that seen in Fig. 9, the normal-state spectral weights are good predictors of the superconducting state's spectral weight. We note that the curves are not symmetrical around $v_0 = 0$, which results from a realistic band structure without particle-hole symmetry.

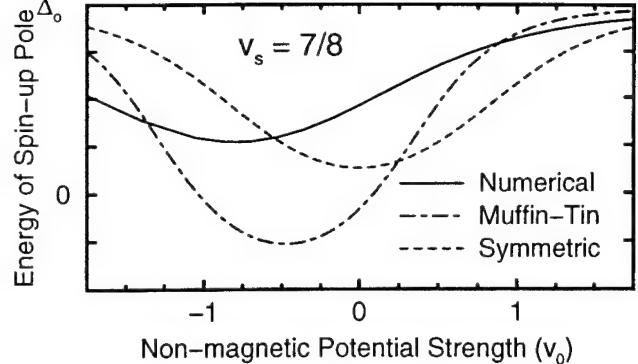


FIG. 14. The energy of the spin-up pole is shown as a function of non-magnetic potential strength for $v_s = 0.875$ and $\xi k_F = 449$. The energy of the spin-down pole is just the negative of the energy of the spin-up pole. The analytic models do not perform particularly well in reproducing the pole energies, although the muffin-tin model does show a similar asymmetry around $v_0 = 0$ to the numerical calculations.

publication.

We now mention an example designed to showcase the drawbacks of Ginzburg-Landau theory. We present in Figure 16 the order parameter for a mixed magnetic and non-magnetic impurity ($v_s = 7/8$, $v_0 = -7/4$, and $\xi k_F = 449$). The order parameter is everywhere larger than in the homogeneous superconductor, however, the presence of a localized state within the gap indicates that superconductivity has been weakened around the impurity. Since Ginzburg-Landau theory focuses on the order parameter and ignores the quasiparticle structure, a Ginzburg-Landau perspective would incorrectly predict an enhancement of superconductivity in the region.

We conclude by attempting to provide some guidance for attempts to extract impurity potentials from STM measurements. Figure 17 shows *normal-state* dI/dV 's for various potentials. These curves should also represent

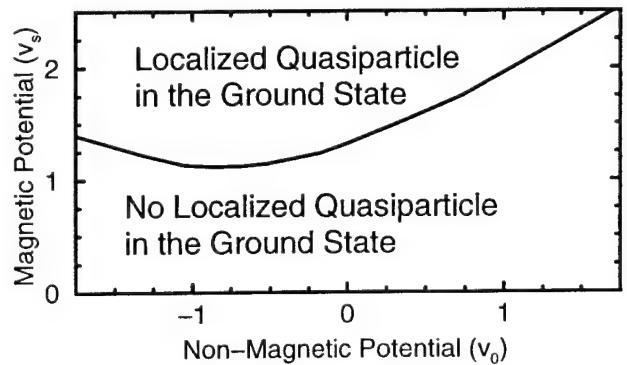


FIG. 15. Calculated boundary between two ground states around the magnetic impurity for $\xi k_F = 449$. For a large enough magnetic impurity strength a quasiparticle is bound in the ground state. The minimum magnetic impurity strength depends on the non-magnetic impurity strength. For still larger magnetic impurity strengths there would be ground states with more than one bound quasiparticle.

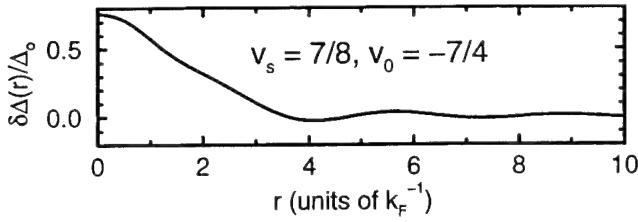


FIG. 16. The change in order parameter around an attractive non-magnetic potential combined with a magnetic potential for $\xi k_F = 449$. The order parameter is larger at the impurity than in the homogeneous superconductor, yet there exists a localized state in the gap.

the frequency-averaged spectral weight measured in the superconducting state (see Eq. (45))⁶⁹.

The enhancement or suppression of spectral weight near the origin is particularly sensitive to v_0 . A measurement of this quantity, the energy of the localized state and the asymmetry of the electron and hole amplitudes at the impurity overconstrains v_0 and v_s , given an assumption of the shape of the potential. To extract information about the potential's detailed shape would require a fitting procedure using the differential conductivity at various positions. If, for some reason, the spin-down amplitude were too small to measure, it may remain possible to constrain the potential strength using the frequency-integrated spectral weight and the localized-state energy.

It seems appropriate to mention again the tendency to normalize spectra according to the LDOS measured at energies much larger than Δ_0 . Since the normal-state LDOS near the impurity changes substantially in the presence of magnetic or non-magnetic impurities, an experiment performed using such a normalization procedure would yield impurity parameters of questionable validity.

D. Pairing Suppression

The pairing potential, $\gamma(\mathbf{x})$ in Eq. (12), may also have spatial structure. When this parameter is changed it induces a change in the order parameter which produces an off-diagonal potential felt by the quasiparticles. We set $v_s = v_0 = 0$ so that there is no magnetic or non-magnetic potential to compete with the order parameter change. Figure 18 shows the order parameter around a short-range suppression,

$$\gamma(\mathbf{x}) = \left[1 - e^{-(k_F r)^2}\right] \gamma_0, \quad (46)$$

for two values of the coherence length. The order parameter is strongly suppressed and since $\gamma(\mathbf{0}) = 0$, $\Delta(\mathbf{0}) = 0$. For long coherence lengths, this change in the order parameter has no effect on the local density of states (shown in Figure 19(a)). It is possible to localize quasiparticle states, however, at shorter coherence lengths. These

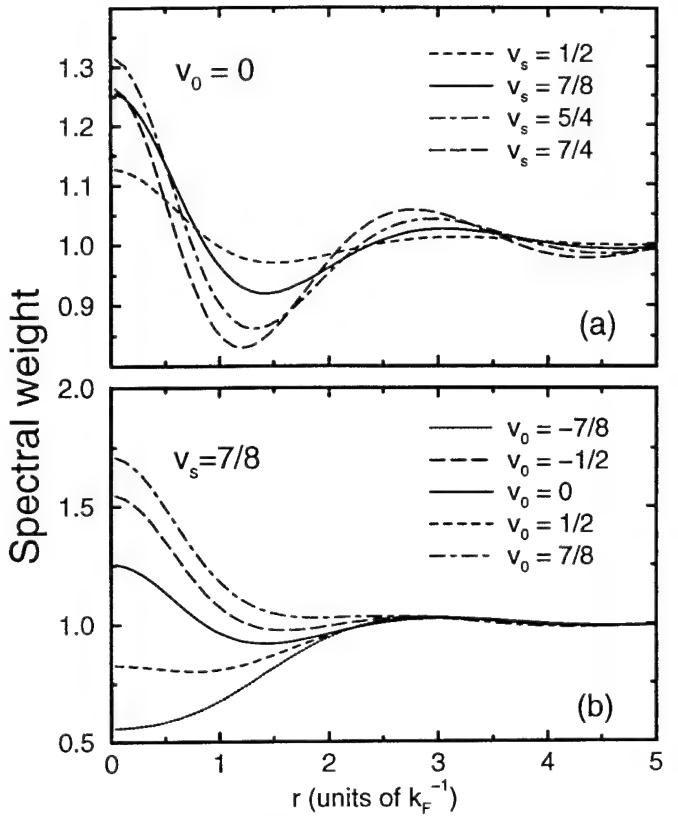


FIG. 17. Spectral weights in the normal state as functions of the distance from the impurity for several combinations of magnetic and non-magnetic potentials (a) only a magnetic potential, and (b) a fixed magnetic potential $v_s = 0.875$ and a varying non-magnetic potential. By making measurements around the impurity in the normal state (or integrating the superconductor's spectrum over a frequency much larger than Δ_0), information about the structure of the impurity may be obtained.

can produce features in the local density of states which are visible. One such case is shown in Figure 19(b). The energy of the localized state is $\omega_0 = (1 - 4 \times 10^{-3})\Delta_0$. Whereas a non-magnetic potential changes the local

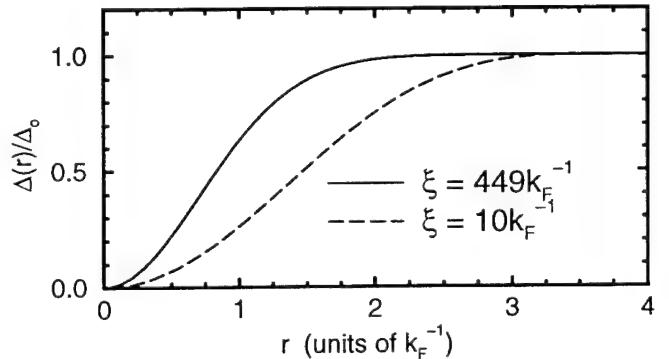


FIG. 18. Order parameter as a function of distance r from a defect with a suppressed pair potential, but no single-particle potential.

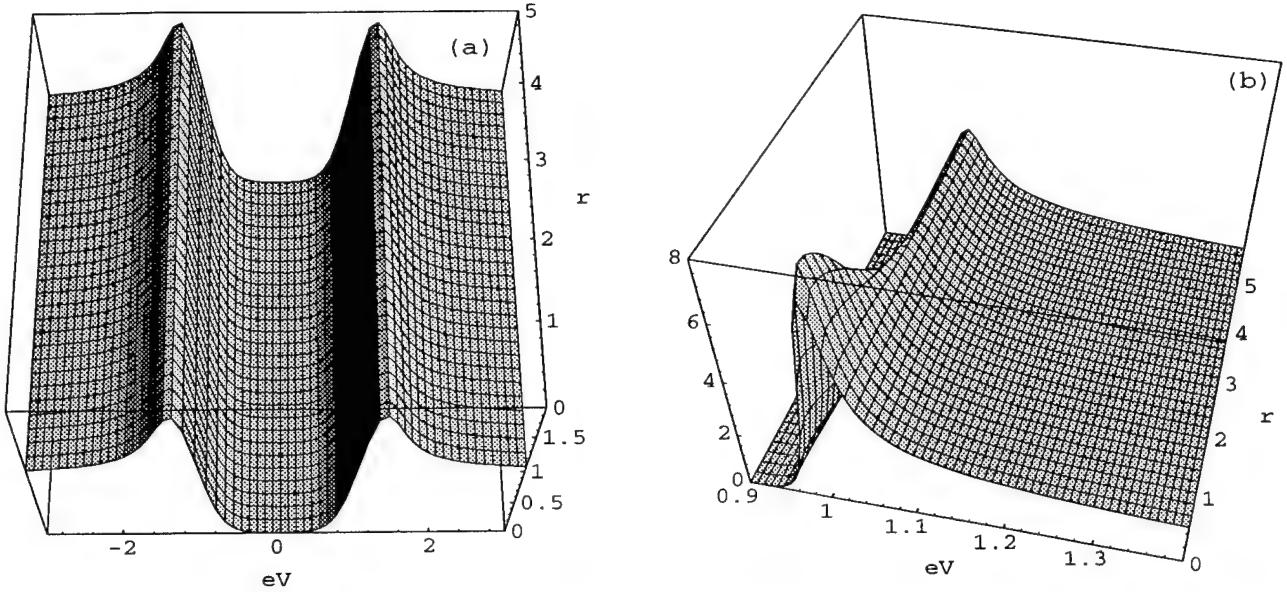


FIG. 19. Differential conductance (LDOS) around a defect with a suppressed pair potential, but no single-particle potential. (a) $\xi k_F = 449$, $k_B T = \Delta_o/7.5$ — there is no evidence of any change in the spectrum due to the order parameter suppression shown in Fig. 18. (b) $\xi k_F = 10$, $k_B T = \Delta_o/100$ — a localized state very near the continuum enhances the continuum edge seen in tunneling near the defect.

density of states without significantly changing the energy gap, a pairing suppression has a very weak effect on both, especially in the long-coherence length limit.

V. STRONG-COUPLING AND ANISOTROPIC ORDER PARAMETERS

A few observations are in order concerning the extension of this formalism to systems where the homogeneous order parameter has important frequency or spatial structure. The Gor'kov equation (Eq. (4)) changes due to the more general form for the off-diagonal potential originating from the order parameter. Taking this opportunity to generalize,

$$\int d\mathbf{x}'' d\mathbf{x}''' [\delta(\mathbf{x} - \mathbf{x}'')\delta(\mathbf{x}'' - \mathbf{x''''}) - \mathbf{g}(\mathbf{x}, \mathbf{x}''; \omega)\mathbf{V}(\mathbf{x}'', \mathbf{x}'''; \omega)] \mathbf{G}(\mathbf{x}''', \mathbf{x}'; \omega) = \mathbf{g}(\mathbf{x}, \mathbf{x}'; \omega) \quad (47)$$

where

$$\mathbf{V}(\mathbf{x}'', \mathbf{x}'''; \omega) = \begin{pmatrix} V_{e\uparrow}(\mathbf{x}''; \omega)\delta(\mathbf{x}'' - \mathbf{x}''') & \delta\Delta(\mathbf{x}'', \mathbf{x}'''; \omega) \\ \delta\Delta(\mathbf{x}'', \mathbf{x}'''; \omega) & V_{h\uparrow}(\mathbf{x}''; \omega)\delta(\mathbf{x}'' - \mathbf{x}''') \end{pmatrix}. \quad (48)$$

The diagonal terms are general potentials, possibly frequency-dependent, effective on spin-up electrons ($V_{e\uparrow}$) and spin-up holes ($V_{h\uparrow}$) for $\omega > 0$. Since this potential is diagonal in frequency, as is the Gor'kov equation, the frequency structure of the order parameter does not add any additional complication to numerically solving the Gor'kov equation. However, the addition of spatial structure to the pairing has added another integral over the volume to the Gor'kov equation, and thus dramatically increased the size of the matrix which needs to be inverted. Fortunately, the range of the order parameter in $|\mathbf{x} - \mathbf{x}'|$ is truncated by the range of the pairing interaction. For the isotropic-gap superconductor we have

considered for most of this paper, the effective pairing interaction is modeled by a delta function in space. It is possible to obtain anisotropic order parameters, including the d -wave order parameters possibly appropriate for high-temperature superconductors, merely by considering pairing with nearest neighbors on a square tight-binding lattice. This model has been implemented with the BdG formalism for non-magnetic impurities³²⁻³⁵ and magnetic impurities³⁸ and vortices⁷⁰ in d -wave superconductors. When formulated on a lattice the addition of nearest-neighbor pairing multiplies the rank of the matrix $\mathbf{M}^{n \rightarrow n}(\omega)$ by $1 + z$, where z is the coordination number of the lattice.

The order parameter's frequency dependence complicates the self-consistency equation (Eq. (12)). It must now be solved for each frequency:

$$\begin{aligned} \Delta(\mathbf{r}, \mathbf{r}'; \omega) Z(\mathbf{r}, \mathbf{r}'; \omega) \\ = \int_{-\infty}^{\infty} d\epsilon n(\epsilon) [\text{Re}F(\mathbf{r}, \mathbf{r}'; \epsilon)] (K_+(\mathbf{r}, \mathbf{r}'; \epsilon, \omega) - U_c) \\ \omega[1 - Z(\mathbf{r}, \mathbf{r}'; \omega)] \\ = \int_{-\infty}^{\infty} d\epsilon n(\epsilon) [\text{Re}G(\mathbf{r}, \mathbf{r}'; \epsilon)] K_-(\mathbf{r}, \mathbf{r}'; \epsilon, \omega), \end{aligned} \quad (49)$$

where K_{\pm} are kernels of the pairing interaction and are different for each mechanism of superconductivity. They can be determined from the homogeneous solution. U_c is a Coulomb factor. Z , the quasiparticle weight, is solved for self-consistently. Incorporating these strong coupling effects allows a determination of the effect of the frequency-dependence of the pairing interaction on the electronic structure around a defect. While selecting the particular location where the STM tip is similar to selecting the momentum of the quasiparticles of interest, selecting the STM voltage indicates which order parameter frequency one wishes to probe.

VI. SUMMARY

The local electronic structure around a defect reflects both the properties of the defect and the medium it is embedded in. For a strong non-magnetic or magnetic impurity in a superconductor, the distortion of the normal-state properties by the strong impurity plays a vital role in the response of the superconducting medium. The LDOS for the inhomogeneous superconductor can be related to the LDOS for the inhomogeneous normal metal via equations like Eq. (42) and Eq. (44). In the case of a non-magnetic impurity with no localized states around it, the LDOS for the inhomogeneous superconductor is merely the normalized LDOS for the inhomogeneous normal metal multiplied by the homogeneous superconductor's density of states. This should suggest some caution for the method used to normalize STM spectra taken at different places on a superconductor's surface.

For the case of a localized state (such as around a magnetic impurity) with angular momentum ℓ , the LDOS for the state is the angular-momentum- ℓ -projected LDOS of

the inhomogeneous normal metal multiplied by a decaying exponential whose range is determined by the energy of the localized state. Since the spin-up band LDOS in the normal state differs from the spin-down band LDOS in the normal state, the electron-like pole of the localized quasiparticle will have different spatial structure than the hole-like pole of the localized quasiparticle.

The self-consistent calculations described here have been performed with a new, powerful, Koster-Slater technique which allows the Gor'kov equation to be solved in principle exactly. Although we have only presented calculations for weak-coupling isotropic order parameters within a free-electron model, we have formulated the extension of the new technique to strong-coupling pairing and general band structures and order parameter symmetries.

ACKNOWLEDGMENTS

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APPENDIX

The expansion of the Green's functions of Eq. (15) suitable for a three-dimensional spherically symmetric situation are detailed here. The homogeneous Green's functions depend on \mathbf{r} and \mathbf{r}' through r , r' , and $\cos \gamma = (\mathbf{r} \cdot \mathbf{r}')/rr'$. Then the Green's functions can be expanded in Legendre polynomials $P_\ell(\cos \gamma)$.

$$g(\mathbf{r}, \mathbf{r}'; \omega) = g(r, r', \cos \gamma; \omega) = \frac{2\ell + 1}{4\pi} \sum_{\ell} g_{\ell}(r, r'; \omega) P_{\ell}(\cos \gamma) \quad (50)$$

and

$$g_{\ell}(r, r'; \omega) = 2\pi \int_{-1}^{1} P_{\ell}(x) g(r, r', x; \omega). \quad (51)$$

Evaluating Eq. (51) for both the normal and anomalous Green's functions of Eq. (15) yields

$$\begin{aligned} g_{\ell}(r, r'; \omega) &= -\frac{\pi^3}{\sqrt{rr'}} \left[i \left(1 + \frac{\omega}{\sqrt{\omega^2 - 1}} \right) J_{\ell+\frac{1}{2}}(\{1 + \sqrt{\omega^2 - 1}/\xi\}r^<) H_{\ell+\frac{1}{2}}^{(1)}(\{1 + \sqrt{\omega^2 - 1}/\xi\}r^>) \right. \\ &\quad \left. - \left(1 - \frac{\omega}{\sqrt{\omega^2 - 1}} \right) J_{\ell+\frac{1}{2}}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^<) H_{\ell+\frac{1}{2}}^{(2)}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^>) \right] \\ f_{\ell}(r, r'; \omega) &= -\frac{i\pi^3}{\sqrt{rr'}} \frac{1}{\sqrt{\omega^2 - 1}} \left[J_{\ell+\frac{1}{2}}(\{1 + \sqrt{\omega^2 - 1}/\xi\}r^<) H_{\ell+\frac{1}{2}}^{(1)}(\{1 + \sqrt{\omega^2 - 1}/\xi\}r^>) \right. \\ &\quad \left. - J_{\ell+\frac{1}{2}}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^<) H_{\ell+\frac{1}{2}}^{(2)}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^>) \right] \end{aligned} \quad (52)$$

$$+ J_{\ell+\frac{1}{2}}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^<) H_{\ell+\frac{1}{2}}^{(2)}(\{1 - \sqrt{\omega^2 - 1}/\xi\}r^>)\Big] \quad (53)$$

where J_ℓ , $H_\ell^{(1)}$, and $H_\ell^{(2)}$ are standard Bessel functions, $r^<$ ($r^>$) is the smaller (larger) of r and r' , ω is in units of Δ_o and r and r' are in units of k_F^{-1} . The Green's functions are in units of N_o .

The Gor'kov equation, Eq. (4), can now be written in a form diagonal in ℓ ,

$$\mathbf{G}_\ell(r, r'; \omega) = \mathbf{g}_\ell(r, r'; \omega) + \int_0^\infty r_n^2 dr_n \mathbf{g}_\ell(r, r_n; \omega) \mathbf{V}(r_n) \mathbf{G}_\ell(r_n, r'; \omega). \quad (54)$$

Thus the three-dimensional integral has been reduced to a one-dimensional radial integral. Since the numerical inversion procedure depends on inverting a matrix whose rank is proportional to the number of spatial points considered, this reduction to a one-dimensional integral dramatically increases the speed of this calculation over a calculation for a three-dimensional potential which is not spherically symmetric.

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